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Form PTO-1390 (Rev. 12-29-99) TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		US DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE
		ATTORNEY'S DOCKET NO C 2037 PCT/US
		U.S. APPLICATION NO. (if known see 37 CFR 1.8) 107049960
INTERNATIONAL APPLICATION NO. PCT/EP00/07849	INTERNATIONAL FILING DATE August 11, 2000	PRIORITY DATE CLAIMED August 20, 1999
TITLE OF INVENTION BRANCHED, SUBSTANTIALLY UNSATURATED FATTY ALCOHOL SULFATES		
APPLICANT(S) FOR DO/EO/US Norbert Huebner, Hans-Christian Raths, Alfred Westfechtel		
Applicant herewith submits to the United States Designated/Elected Office (EO/DO/US) the following items and other information:		
<p>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.</p> <p>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.</p> <p>3. <input type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39 (1).</p> <p>4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.</p> <p>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)).</p> <p>a. <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau).</p> <p>b. <input checked="" type="checkbox"/> has been transmitted by the International Bureau.</p> <p>c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</p> <p>6. <input checked="" type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)).</p> <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))</p> <p>a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau).</p> <p>b. <input type="checkbox"/> have been transmitted by the International Bureau.</p> <p>c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</p> <p>d. <input checked="" type="checkbox"/> have not been made and will not be made.</p> <p>8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</p> <p>9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). (UNEXECUTED)</p> <p>10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</p>		
Items 11. to 16. below concern other document(s) or information included:		
<p>11. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</p> <p>12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</p> <p>13. <input checked="" type="checkbox"/> A FIRST preliminary amendment</p> <p><input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>14. <input type="checkbox"/> A substitute specification.</p> <p>15. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>16. <input type="checkbox"/> Other items or information:</p>		
<p>"Express Mail Post Office to Addressee" service Mailing Label Number <u>EL541614378US</u></p>		

U.S. Application No. (if known, see 37 CFR 1.5)	INTERNATIONAL APPLICATION NO.	ATTORNEY'S DOCKET NUMBER	
107049960	PCT/EP00/07849	C 2037 PCT/US	
<p>17. <input checked="" type="checkbox"/> The following fees are submitted:</p> <p>BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5)):</p> <p>Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO.....\$1,040.00</p> <p>International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO.....\$890.00</p> <p>International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International search fee (37 CFR 1.445(a)(2)) paid to USPTO\$740.00</p> <p>International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$710.00</p> <p>International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4).....\$100.00</p>		<p>CALCULATIONS PTO USE ONLY</p>	
<p>ENTER APPROPRIATE BASIC FEE AMOUNT = \$ 890</p> <p>Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).</p>		<p>\$ 0</p>	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total Claims	9 - 20 =	0	0 X \$18.00
Independent Claims	3 - 3 =	0	0 X \$84.00
Multiple dependent claims (s)(if applicable)	0	+ \$280.00	\$ 0
TOTAL OF ABOVE CALCULATIONS =		\$ 890	
Reduction of $\frac{1}{2}$ for filing by small entity, if applicable. A Small Entity Statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).		\$ 0	
SUBTOTAL =		\$ 890	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$ 0	
TOTAL NATIONAL FEE =		\$ 890	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property		\$ 0	
TOTAL FEES ENCLOSED =		\$ 890	
		Amount to be: refunded:	\$ -----
		charged:	\$ 890.00
<p>a. <input type="checkbox"/> A check in the amount of \$ _____ to cover the above fees is enclosed.</p> <p>b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>50-1177</u> in the amount of <u>\$890.00</u> to cover the above fees. A triplicate copy of this sheet is enclosed. Order No. <u>02-0080</u>.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>50-1177</u>. A triplicate copy of this sheet is enclosed.</p> <p>NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.</p>			
<p>SEND ALL CORRESPONDENCE TO: Customer Label No. 23657</p> <p> _____ SIGNATURE</p> <p>John E. Drach NAME ATTORNEY FOR APPLICANT 32,891 REGISTRATION NUMBER</p>			

"Express Mail" mailing label number EL541614378US.

PATENT
Docket No. C 2037 PCT/US

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

RE: PCT/EP00/07849
International Filing Date: August 11, 2000
Priority Date Claimed: August 20, 1999
Applicant: Huebner, et al.
Title: BRANCHED, SUBSTANTIALLY UNSATURATED FATTY
ALCOHOL SULFATES
Applicants' Reference: C 2037 PCT/US

PRELIMINARY AMENDMENT

Commissioner for Patents
Box PCT
Washington, DC 20231

ATTN: DO/EO/US

Sir:

Before examination, in the national stage for the United States, of the above-captioned application under the Patent Convention Treaty, please amend as follows the translation supplied herewith of the application:

In the Specification:

Please delete all text above line 8, of page 1, and replace the deleted matter with the following new section headings and new paragraph:

—TITLE OF THE INVENTION

Branched Substantially Unsaturated Fatty Alcohol Sulfates

BACKGROUND OF THE INVENTION

This invention relates generally to anionic surfactants and, more particularly, to substantially unsaturated fatty alcohol sulfates which, through the presence of branches in the hydrocarbon chain, are distinguished from linear homologs by significantly improved properties, to a process for their production and to their use for the production of surface-active compositions.—

Preliminary Amendment of U.S. National Stage for International Application
PCT/EP00/07849 filed on August 11, 2000

On a separate, new page 46, following page 45, please add the following new header and paragraph containing an Abstract of the Disclosure:

--ABSTRACT OF THE DISCLOSURE

Branched, substantially unsaturated fatty alcohol sulfates are produced by a process which comprises the steps of: (a) dimerizing unsaturated C₁₆₋₂₂ fatty acid to form a dimer fraction and a monomer fraction comprised of branched, substantially unsaturated fatty acids and straight chain saturated fatty acids, (b) removing the monomer fraction from the dimerization step, (c) converting the branched, substantially unsaturated fatty acids from step (b) into the corresponding fatty acid methyl esters, (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and (e) sulfating and neutralizing the branched, substantially unsaturated fatty alcohols. The fatty alcohol sulfates thus produced exhibit improved performance properties and greater oxidative stability than standard unsaturated fatty alcohol sulfates.--

In the claims:

Please cancel claims 1-10.

Please add the following new claims 11-19.

11. (New) A branched, substantially unsaturated fatty alcohol sulfate which is the product of the process comprising the steps of: (a) dimerizing an unsaturated C₁₆₋₂₂ fatty acid to form a dimer fraction and a monomer fraction comprised of branched, substantially unsaturated fatty acids and straight chain saturated fatty acids, (b) separating the monomer fraction from the dimer fraction, (c) converting the branched, substantially unsaturated fatty acids in the monomer fraction into the corresponding fatty acid methyl esters, (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and (e) sulfating and neutralizing the branched, substantially unsaturated fatty alcohols.

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PCT/EP00/07849 filed on August 11, 2000**

12. (New) A process for the production of branched, substantially unsaturated fatty alcohol ether sulfates comprising the steps of: (a) dimerizing an unsaturated C₁₆₋₂₂ fatty acid to form a dimer fraction and a monomer fraction comprised of branched, substantially unsaturated fatty acids and straight chain saturated fatty acids, (b) separating the monomer fraction from the dimer fraction, (c) converting the branched, substantially unsaturated fatty acids in the monomer fraction into the corresponding fatty acid methyl esters, (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and (e) sulfating and neutralizing the branched, substantially unsaturated fatty alcohols.

13. (New) The process of claim 12 wherein the branched, substantially unsaturated fatty alcohols are sulfated with sulfur trioxide or chlorosulfonic acid.

14. (New) The process of claim 12 wherein step (e) is carried out in a falling-film reactor.

15. (New) The process of claim 12 wherein step (e) is carried out at temperature of from about 25 to about 90°C.

16. (New) The process of claim 12 wherein in step (e) the molar ratio of fatty alcohol to sulfating agent is from about 1:0.95 to about 1:1.8.

17. (New) A process for the production of branched, substantially unsaturated fatty alcohol ether sulfates comprising the steps of: (a) dimerizing an unsaturated C₁₆₋₂₂ fatty acid to form a dimer fraction and a monomer fraction comprised of branched, substantially unsaturated fatty acids and straight chain saturated fatty acids, (b) removing the straight chain saturated fatty acids from the monomer fraction by fractional crystallization and optionally separating the crystallization solvent by distillation, (c) converting the branched, substantially unsaturated fatty

**Preliminary Amendment of U.S. National Stage for International Application
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acids in the monomer fraction into the corresponding fatty acid methyl esters, (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and (e) sulfating and neutralizing the branched, substantially unsaturated fatty alcohols.

18. (New) The process of claim 12 wherein the methyl esters and/or the corresponding branched, substantially unsaturated fatty alcohols are distilled prior to step (e).
19. (New) The process of claim 17 wherein the methyl esters and/or the corresponding branched, substantially unsaturated fatty alcohols are distilled prior to step (e).

**Preliminary Amendment of U.S. National Stage for International Application
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REMARKS

Claims 11-19 are currently pending in the instant application.

The Specification has been amended to include the preferred section headings pursuant to 37 C.F.R. §1.77. It is submitted that the amendments to the Specification made herein introduce no new matter. Their entry is therefore proper and respectfully requested. An Abstract of the Disclosure has been added on a separate sheet following the claims.

Original claims 1-10 have been canceled and replaced with new claims 11-19 solely for the purpose of improving clarity and grammar, which may suffer in translation, and not for any reason related to the statutory requirements for a patent. New claims 11-19 have not been added in response to any rejection, or in anticipation of any rejection related to the statutory requirements for a patent. Applicants respectfully submit that the scope of new claims 11-19 corresponds to the scope of original claims 1-10 and that new claims 11-19 are no narrower than original claims 1-10. Furthermore, although a moot point in view of their cancellation, Applicants respectfully submit that original claims 1-10 satisfied the requirements of 35 U.S.C. §112, as filed. New claims 11-19 are supported by the specification and no new matter has been introduced. Entry is therefore proper and respectfully requested. Prompt examination of the instant application in view of the amendments made herein is respectfully requested.

Respectfully submitted,



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ABSTRACT OF THE DISCLOSURE

Branched, substantially unsaturated fatty alcohol sulfates are produced by a process which comprises the steps of: (a) dimerizing unsaturated C₁₆₋₂₂ fatty acid to form a dimer fraction and a monomer fraction comprised of branched, substantially unsaturated fatty acids and straight chain saturated fatty acids, (b) removing the monomer fraction from the dimerization step, (c) converting the branched, substantially unsaturated fatty acids from step (b) into the corresponding fatty acid methyl esters, (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and (e) sulfating and neutralizing the branched, substantially unsaturated fatty alcohols. The fatty alcohol sulfates thus produced exhibit improved performance properties and greater oxidative stability than standard unsaturated fatty alcohol sulfates.

Branched Substantially Unsaturated Fatty Alcohol Sulfates

Field of the Invention

This invention relates generally to anionic surfactants and, more particularly, to substantially unsaturated fatty alcohol sulfates which, through the presence of branches in the hydrocarbon chain, are distinguished from linear homologs by significantly improved properties, to

- 5 a process for their production and to their use for the production of surface-active compositions.

Prior Art

Sulfates of unsaturated fatty alcohols, which are essentially obtained
10 by sulfation and subsequent neutralization of the corresponding tallow-based alkenols, are important raw materials for the production of both cosmetic preparations and laundry detergents, dishwashing detergents and cleaning products. The favorable properties of these substances are linked to the presence of the double bond in the molecule which also presents
15 problems because the unsaturated fatty alcohol ether sulfates readily fall victim to autoxidation which is associated with discoloration and unwanted chemical changes (for example formation of peroxides and hydroperoxides).

Accordingly, it is clear that there is a need on the market for
20 unsaturated fatty alcohol sulfates with improved oxidation stability or suitable substitutes which possess at least equivalent performance properties. However, more or less pure isostearyl alcohol sulfates have hitherto been the only alternatives to unsaturated fatty alcohol sulfates. To produce these more or less pure isostearyl alcohol sulfates, however, oleic acid first has to be dimerized, the fraction of monomeric branched fatty acids separated off, hydrogenated and subjected to fractional

crystallization, the liquid fraction accumulating, which is rich in isostearic acid, has to be removed and esterified with methanol and the esters obtained subsequently hydrogenated to form the alcohols which, finally, are converted into the sulfates.

- 5 The process described above is technically complicated by the two hydrogenation steps and, in the isostearyl alcohol sulfates, provides substitutes which can only replace the unsaturated fatty alcohol sulfates to a limited extent. Accordingly, the problem addressed by the present invention was to provide unsaturated unsaturated fatty alcohol sulfates
10 which would be distinguished by improved performance properties and preferably by higher oxidation stability.

Description of the Invention

The present invention relates to branched, substantially unsaturated fatty alcohol sulfates which are obtainable by

- 15 (a) dimerizing unsaturated C₁₆₋₂₂ fatty acids in known manner,
 (b) removing the monomer fraction accumulating in the dimerization step,
 (c) converting the branched, substantially unsaturated fatty acids present in
20 this fraction into the corresponding fatty acid methyl esters,
 (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and
 (e) sulfating and neutralizing the branched, substantially unsaturated fatty
25 alcohols in known manner.

It has surprisingly been found that the branched, substantially unsaturated fatty alcohol sulfates have distinctly improved autoxidation stability compared with linear homologs having the same chain length and
30 the same iodine value. Further advantages include improved wetting

behavior, quicker solubility in cold water and easier biodegradability.

The present invention also relates to a process for the production of branched, substantially unsaturated fatty alcohol sulfates in which

- 5 (a) unsaturated C₁₆₋₂₂ fatty acids are dimerized in known manner,
- (b) the monomer fraction accumulating in the dimerization step is removed,
- (c) the branched, substantially unsaturated fatty acids present in this fraction are converted into the corresponding fatty acid methyl esters,
- 10 (d) the branched, substantially unsaturated fatty acid methyl esters are hydrogenated with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols which are then
- (e) sulfated and neutralized in known manner.

15 Production of the fatty alcohols

The dimerization of fatty acids and the recovery of monomer fatty acids from the dimers is sufficiently well-known from the prior art, cf. for example the overviews by A. Behr et al. [*Fat Sci. Technol.* 93, 340 (1991)] and by H. Möhring et al. [*ibid.* 94, 41 (1992) and 94, 241 (1992)]. The sequence of steps (a) to (d) gives branched, substantially unsaturated fatty alcohols with iodine values of 45 to 85 on the basis of dimerized, preferably monounsaturated C₁₆₋₂₂ fatty acids, i.e. oleic acid, elaidic acid, petroselinic acid, gadoleic acid and erucic acid and mixtures thereof. This is without doubt entirely adequate for a number of applications. However, in cases where fatty compounds with a relatively high content of unsaturated compounds are required, it is advisable to subject the monomer fraction accumulating in the dimerization step to fractional crystallization and then to subject the liquid phase obtained to esterification, optionally after distillation. The fatty acid obtained and its methyl esters represent an already fairly pure isooleic acid or isooleic acid methyl ester with iodine

values of 75 to 95. In any event, it is advisable to subject the methyl esters and/or the fatty alcohols to distillation and/or fractional crystallization ("winterizing"). The esterification of the fatty acids with methanol is carried out by known methods and is intended to produce methyl esters which are

5 comparatively easy to hydrogenate. Instead of the methyl esters, other lower alkyl esters, for example ethyl, propyl or butyl esters, may of course also be produced and subsequently hydrogenated. The choice of the alcohol is basically not critical and is solely determined by economic criteria and availability. Instead of the methyl or lower alkyl esters, it is also

10 possible in principle directly to esterify the fatty acids, although this does involve the use of special catalysts which do not form salts with the acids. In addition, the reactor material has to be corrosion-resistant. The hydrogenation of the unsaturated methyl esters to form the corresponding alcohols may also be carried out in known manner. Corresponding

15 processes and catalysts, particularly those based on copper and zinc, are disclosed for example in the following documents: DE 43 357 81 C1, EP 0 602 108 B1, US 3,193,586 and US 3,729,520 (Henkel); reference is expressly made to the disclosures of these documents.

20 Sulfation and neutralization

The conversion of the fatty alcohols previously obtained into the sulfates may also be carried out in known manner. The attack of the sulfating agent may be directed both at the hydroxyl group and at the double bond. However, since the sulfation process takes place about 10

25 times more quickly than sulfonation, i.e. the addition of sulfur trioxide onto the double bond, at low temperatures of the order of 30°C, sulfates are predominantly obtained (i.e. to an extent of more than 90% by weight). The reaction of the branched, substantially unsaturated fatty alcohols, for example with gaseous sulfur trioxide, may be carried out in the same way

30 as described for fatty acid lower alkyl esters in J. Falbe (ed.), "Surfactants

in Consumer Products"; Springer Verlag, Berlin-Heidelberg, 1987, page 61, reactors operating on the falling-film principle being preferred. In this known process, the sulfur trioxide is diluted with an inert gas, preferably air or nitrogen, and used in the form of a gas mixture which

5 contains the sulfonating agent in a concentration of 1 to 8% by volume and, more particularly, 2 to 5% by volume. The molar ratio of fatty alcohol to sulfating agent is 1:0.95 to 1:1.8, preferably from 1:1.0 to 1:1.6 and more preferably from 1:1.3 to 1:1.5. The sulfation reaction is generally carried out at temperatures of 25 to 90°C and preferably at temperatures of 35 to

10 80°C. Instead of sulfur trioxide, chlorosulfonic acid or amidosulfonic acid may also be used as the sulfonating agent. The acidic sulfates accumulating during the reaction are stirred into aqueous bases, neutralized and adjusted to a pH value of 6.5 to 8.5. The neutralization step is carried out with bases selected from the group consisting of alkali

15 metal hydroxides, such as sodium, potassium and lithium hydroxide, alkaline earth metal oxides and hydroxides, such as magnesium oxide, magnesium hydroxide, calcium oxide and calcium hydroxide, ammonia, mono-, di- and tri-C₂₋₄-alkanolamines, for example mono-, di- and triethanolamine, and primary, secondary or tertiary C₁₋₄ alkyl amines. The

20 neutralization bases are preferably used in the form of 5 to 55% by weight aqueous solutions, 25 to 50% by weight aqueous sodium hydroxide solution being preferred. After neutralization, the sulfates obtainable by the process according to the invention are present as aqueous solutions with an active substance content of 20 to 80% by weight and preferably 30 to

25 50% by weight. After neutralization, the sulfates may be bleached in known manner by addition of hydrogen peroxide or sodium hypochlorite solution to achieve a further lightening in color desirable for many applications. 0.2 to 2% by weight of hydrogen peroxide – expressed as 100% by weight substance – or corresponding quantities of sodium

30 hypochlorite, based on the solids content in the solution of the sulfates, are

used for this purpose. The pH value of the solutions may be kept constant using suitable buffers, for example sodium phosphate or citric acid. In addition, preservation, for example with formaldehyde solution, p-hydroxybenzoate, sorbic acid or other known preservatives, is advisable for 5 stabilization against bacterial infestation.

Commercial Applications

The new branched substantially unsaturated fatty alcohol sulfates are distinguished by particular stability to oxidation and are therefore 10 suitable for the production of surface-active compositions, preferably laundry detergents, dishwashing detergents, cleaners and softeners, and cosmetic and/or pharmaceutical preparations in which they may be present in quantities of 1 to 50% by weight, preferably 5 to 35% by weight and more preferably 10 to 25% by weight.

15

Laundry detergents, dishwashing detergents, cleaners and softeners

Where the branched, substantially unsaturated fatty alcohol sulfates according to the invention are used as raw materials for the production of laundry detergents, dishwashing detergents, cleaners or softeners, they are 20 normally present in liquid form, i.e. as aqueous solutions or pastes; for the production of detergent powders, the water-containing mixtures may be subsequently dried. Liquid preparations may have a non-aqueous component of 5 to 50% by weight and preferably 15 to 35% by weight. In the most simple case, they are aqueous solutions of the mixtures 25 mentioned. However, the liquid detergents may also be substantially water-free compositions. "Substantially water-free" in the context of the present invention means that the composition preferably contains no free water which is not bound as water of crystallization or in a comparable form. In some cases, small quantities of free water are tolerable, more 30 particularly quantities of up to 5% by weight. The compositions used in the

detergent field may contain other typical ingredients such as, for example, builders, bleaching agents, bleach activators, solvents, detergency boosters, enzymes, enzyme stabilizers, viscosity regulators, redeposition inhibitors, optical brighteners, soil repellents, foam inhibitors, inorganic salts

5 and fragrances and dyes.

Suitable **liquid builders** are ethylenediamine tetraacetic acid, nitrilotriacetic acid, citric acid and inorganic phosphonic acids such as, for example, the neutrally reacting sodium salts of 1-hydroxyethane-1,1-diphosphonate which may be present in quantities of 0.5 to 5% by weight

10 and preferably 1 to 2% by weight.

A suitable **solid builder** is, in particular, finely crystalline zeolite containing synthetic and bound water, such as detergent-quality zeolite NaA. However, zeolite NaX and mixtures of NaA and NaX are also suitable. The zeolite may be used in the form of a spray-dried powder or

15 even as an undried stabilized suspension still moist from its production. Where the zeolite is used in the form of a suspension, the suspension may contain small additions of nonionic surfactants as stabilizers, for example 1 to 3% by weight – based on zeolite – of ethoxylated C₁₂₋₁₈ fatty alcohols containing 2 to 5 ethylene oxide groups or ethoxylated isotridecanols.

20 Suitable zeolites have a mean particle size of less than 10 µm (volume distribution, as measured by the Coulter Counter Method) and contain preferably 18 to 22% by weight and more preferably 20 to 22% by weight of bound water. Suitable substitutes or partial substitutes for zeolites are crystalline layer-form sodium silicates with the general formula

25 NaMSi_xO_{2x+1}yH₂O, where M is sodium or hydrogen, x is a number of 1.9 to 4 and y is a number of 0 to 20, preferred values for x being 2, 3 or 4. Crystalline layer silicates such as these are described, for example, in European patent application EP 0 164 514 A. Preferred crystalline layer silicates are those in which M in the general formula stands for sodium and

30 x assumes the value 2 or 3. Both β- and γ-sodium disilicates

$\text{Na}_2\text{Si}_2\text{O}_5\text{yH}_2\text{O}$ are particularly preferred, β -sodium disilicate being obtainable for example by the process described in International patent application WO 91/08171. Powder-form detergents based on the branched, substantially unsaturated fatty alcohol sulfates according to the

5 invention preferably contain 10 to 60% by weight of zeolite and/or crystalline layer silicates as solid builders, mixtures of zeolite and crystalline layer silicates in any ratio being particularly advantageous. In one particularly preferred embodiment, the detergents contain 20 to 50% by weight of zeolite and/or crystalline layer silicates. Particularly preferred

10 detergents contain up to 40% by weight of zeolite and, more particularly, up to 35% by weight of zeolite, based on water-free active substance. Other suitable ingredients of the detergents are water-soluble amorphous silicates which are preferably used in combination with zeolite and/or crystalline layer silicates. Particularly preferred detergents are those which

15 contain above all sodium silicate with a molar ratio of Na_2O to SiO_2 (modulus) of 1:1 to 1:4.5 and preferably 1:2 to 1:3.5. The amorphous sodium silicate content of the detergents is preferably up to 15% by weight and more preferably from 2 to 8% by weight. Phosphates, such as tripolyphosphates, pyrophosphates and orthophosphates, may also be

20 present in the detergents in small quantities. The phosphate content of the detergents is preferably up to 15% by weight and, more particularly, from 0 to 10% by weight. In addition, the detergents may contain layer silicates of natural and synthetic origin. Corresponding layer silicates are known, for example, from patent applications DE 23 34 899 B, EP 0 026 529 A and

25 DE 35 26 405 A. Their suitability for use is not confined to a particular composition or structural formula. However, smectites are preferred, bentonites being particularly preferred. Suitable layer silicates which belong to the group of water-swellable smectites are, for example, those corresponding to the following general formulae:

WO 01/14325

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$(OH)_4Si_{8-y}Al_y(Mg_xAl_{4-x})O_{20}$	montmorillonite
$(OH)_4Si_{8-y}Al_y(Mg_{6-z}Li_z)O_{20}$	hectorite
$(OH)_4Si_{8-y}Al_y(Mg_{6-z}Al_z)O_{20}$	saponite

- 5 where x = 0 to 4, y = 0 to 2 and z = 0 to 6. In addition, small quantities of iron may be incorporated in the crystal lattice of the layer silicates corresponding to the above formulae. By virtue of their ion-exchanging properties, the layer silicates may also contain hydrogen, alkali metal and alkaline earth metal ions, more particularly Na^+ and Ca^{2+} . The quantity of
- 10 water of hydration is generally in the range from 8 to 20% by weight and is dependent upon the degree of swelling and upon the processing method. Suitable layer silicates are known, for example, from **US 3,966,629**, **US 4,062, 647**, **EP 0 026 529 A** and **EP 0 028 432 A**. Layer silicates which have been substantially freed from calcium ions and strongly coloring iron
- 15 ions by an alkali treatment are preferably used. Useful organic builders are, for example, the polycarboxylic acids preferably used in the form of their sodium salts, such as citric acid, adipic acid, succinic acid, glutaric acid, tartaric acid, sugar acids, aminocarboxylic acids, nitrilotriacetic acid (NTA), providing their use is not ecologically unsafe, and
- 20 mixtures thereof. Preferred salts are the salts of polycarboxylic acids, such as citric acid, adipic acid, succinic acid, glutaric acid, tartaric acid, sugar acids and mixtures thereof. Suitable polymeric polycarboxylates are, for example, the sodium salts of polyacrylic acid or polymethacrylic acid, for example those with a relative molecular weight of 800 to 150,000 (based
- 25 on acid). Suitable copolymeric polycarboxylates are, in particular, those of acrylic acid with methacrylic acid and acrylic acid or methacrylic acid with maleic acid. Copolymers of acrylic acid with maleic acid which contain 50 to 90% by weight of acrylic acid and 50 to 10% by weight of maleic acid are particularly suitable. Their relative molecular weight, based on free acids,
- 30 is generally in the range from 5,000 to 200,000, preferably in the range

from 10,000 to 120,000 and more preferably in the range from 50,000 to 100,000. It is not absolutely essential to use polymeric polycarboxylates. However, if polymeric polycarboxylates are used, detergents containing biodegradable polymers, for example terpolymers which contain acrylic acid and maleic acid or salts thereof and vinyl alcohol or vinyl alcohol derivatives as monomers or acrylic acid and 2-alkyl allyl sulfonic acid or salts thereof and sugar derivatives as monomers are preferred. The terpolymers obtained in accordance with the teaching of German patent applications **DE 42 21 381 A** and **DE 43 00 772 A** are particularly preferred. Other suitable builders are polyacetals which may be obtained by reacting dialdehydes with polyol carboxylic acids containing 5 to 7 carbon atoms and at least 3 hydroxyl groups, for example as described in European patent application **EP 0 280 223 A**. Preferred polyacetals are obtained from dialdehydes, such as glyoxal, glutaraldehyde, terephthalaldehyde and mixtures thereof and from polyol carboxylic acids, such as gluconic acid and/or glucoheptonic acid.

Among the compounds yielding hydrogen peroxide in water which are used as **bleaching agents**, sodium perborate tetrahydrate and sodium perborate monohydrate are particularly important. Other suitable bleaching agents are, for example, peroxycarbonate, citrate perhydrates and salts of peracids, such as perbenzoates, peroxyphthalates or diperoxydodecanoic acid. They are normally used in quantities of 8 to 25% by weight. Sodium perborate monohydrate is preferred and is used in quantities of 10 to 20% by weight and preferably in quantities of 10 to 15% by weight. By virtue of its ability to bind free water to form the tetrahydrate, it contributes towards increasing the stability of the detergent.

In order to obtain an improved bleaching effect where washing is carried out at temperatures of 60°C or lower, **bleach activators** may be incorporated in the preparations. Examples of bleach activators are N-acyl and O-acyl compounds which form organic peracids with hydrogen per-

- oxide, preferably N,N'-tetraacylated diamines, also carboxylic anhydrides and esters of polyols, such as glucose pentaacetate. The bleach activator content of bleach-containing detergents is in the usual range, i.e. preferably between 1 and 10% by weight and more preferably between 3 and 8% by weight. Particularly preferred bleach activators are N,N,N',N'-tetraacetyl ethylenediamine and 1,5-diacetyl-2,4-dioxohexahydro-1,3,5-triazine.

Suitable organic **solvents** are, for example, monohydric and/or polyhydric alcohols containing 1 to 6 carbon atoms and preferably 1 to 4 carbon atoms. Preferred alcohols are ethanol, propane-1,2-diol, glycerol and mixtures thereof. The detergents preferably contain 2 to 20% by weight and more preferably 5 to 15% by weight of ethanol or a mixture of ethanol and propane-1,2-diol or, more particularly, of ethanol and glycerol. In another possible embodiment, the preparations contain polyethylene glycol with a relative molecular weight of 200 to 2,000 and preferably up to 15 600 in quantities of 2 to 17% by weight either in addition to the monohydric and/or polyhydric alcohols containing 1 to 6 carbon atoms or on its own. Suitable hydrotropes are, for example, toluene sulfonate, xylene sulfonate, cumene sulfonate or mixtures thereof.

Suitable **enzymes** are those from the class of proteases, lipases, 20 amylases, cellulases and mixtures thereof. Enzymes obtained from bacterial strains or fungi, such as *Bacillus subtilis*, *Bacillus licheniformis* and *Streptomyces griseus*, are particularly suitable. Proteases of the subtilisin type are preferably used, proteases obtained from *Bacillus lentus* being particularly preferred. They may be used in quantities of about 0.2 to 25 about 2% by weight. The enzymes may be adsorbed onto supports and/or embedded in membrane materials to protect them against premature decomposition. In addition to the monohydric and polyhydric alcohols and the phosphonates, the detergents may contain other enzyme stabilizers. For example, 0.5 to 1% by weight of sodium formate may be used. It is 30 also possible to use proteases which are stabilized with soluble calcium

salts and which have a calcium content of preferably about 1.2% by weight, based on the enzyme. However, it is of particular advantage to use boron compounds, for example boric acid, boron oxide, borax and other alkali metal borates, such as the salts of orthoboric acid (H_3BO_3), metaboric acid

5 (HBO_2) and pyroboric acid (tetraboric acid $H_2B_4O_7$).

Suitable **viscosity regulators** are, for example, hydrogenated castor oil, salts of long-chain fatty acids, which are preferably used in quantities of 0 to 5% by weight and more preferably in quantities of 0.5 to 2% by weight, for example sodium, potassium, aluminium, magnesium and 10 titanium stearates or the sodium and/or potassium salts of behenic acid, and other polymeric compounds. Preferred other polymeric compounds include polyvinyl pyrrolidone, urethanes and the salts of polymeric polycarboxylates, for example homopolymeric or copolymeric polyacrylates, polymethacrylates and, in particular, copolymers of acrylic acid with maleic 15 acid, preferably those of 50% to 10% maleic acid. The relative molecular weight of the homopolymers is generally between 1,000 and 100,000 while the relative molecular weight of the copolymers is between 2,000 and 200,000 and preferably between 50,000 and 120,000, based on the free acid. Water-soluble polyacrylates which are crosslinked, for example, with 20 about 1% of a polyallyl ether of sucrose and which have a relative molecular weight above 1,000,000 are also particularly suitable. Examples include the polymers with a thickening effect obtainable under the name of Carbopol® 940 and 941. The crosslinked polyacrylates are preferably used in quantities of not more than 1% by weight and more preferably in 25 quantities of 0.2 to 0.7% by weight. The detergents may additionally contain about 5 to 20% by weight of a partly esterified copolymer of the type described in European patent application EP 0 367 049 A. These partly esterified polymers are obtained by copolymerization of (a) at least one C_{4-28} olefin or mixtures of at least one C_{4-28} olefin with up to 20 mole-% 30 of C_{1-28} alkyl vinyl ethers and (b) ethylenically unsaturated dicarboxylic

- anhydrides containing 4 to 8 carbon atoms in a molar ratio of 1:1 to form copolymers with K values of 6 to 100 and subsequent partial esterification of the copolymers with reaction products, such as C₁₋₁₃ alcohols, C₈₋₂₂ fatty acids, C₁₋₁₂ alkyl phenols, secondary C₂₋₃₀ amines or mixtures thereof, with
- 5 at least one C₂₋₄ alkylene oxide or tetrahydrofuran and hydrolysis of the anhydride groups of the copolymers to carboxyl groups, the partial esterification of the copolymers being continued to such an extent that 5 to 50% of the carboxyl groups of the copolymers are esterified. Preferred copolymers contain maleic anhydride as the ethylenically unsaturated
- 10 dicarboxylic anhydride. The partly esterified copolymers may be present either in the form of the free acid or preferably in partly or completely neutralized form. The copolymers are advantageously used in the form of an aqueous solution, more particularly in the form of a 40 to 50% by weight solution. The copolymers not only contribute towards the single wash cycle
- 15 and multiple wash cycle performance of the liquid detergent, they also promote a desirable reduction in viscosity of the concentrated liquid detergents. By using these partly esterified copolymers, it is possible to obtain concentrated aqueous liquid detergents which flow under the sole effect of gravity, i.e. without any need for other shear forces. In a preferred
- 20 embodiment, the concentrated aqueous liquid detergents contain partly esterified copolymers in quantities of 5 to 15% by weight and, more particularly, in quantities of 8 to 12% by weight.

The function of **redeposition inhibitors** is to keep the soil detached from the fibers suspended in the wash liquor and thus to prevent

25 discoloration. Suitable redeposition inhibitors are water-soluble, generally organic colloids, for example the water-soluble salts of polymeric carboxylic acids, glue, gelatin, salts of ether carboxylic acids or ether sulfonic acids of starch or cellulose or salts of acidic sulfuric acid esters of cellulose or starch. Water-soluble polyamides containing acidic groups are also

30 suitable for this purpose. Soluble starch preparations and other starch

products than those mentioned above, for example degraded starch, aldehyde starches, etc., may also be used. Polyvinyl pyrrolidone is also suitable. However, cellulose ethers, such as carboxymethyl cellulose, methyl cellulose, hydroxyalkyl cellulose, and mixed ethers, such as methyl 5 hydroxyethyl cellulose, methyl hydroxypropyl cellulose, methyl carboxymethyl cellulose and mixtures thereof, and polyvinyl pyrrolidone are preferably used, for example in quantities of 0.1 to 5% by weight, based on the detergent.

The detergents may contain derivatives of diaminostilbene disulfonic 10 acid or alkali metal salts thereof as **optical brighteners**. Suitable optical brighteners are, for example, salts of 4,4'-bis-(2-anilino-4-morpholino-1,3,5-triazinyl-6-amino)-stilbene-2,2'-disulfonic acid or compounds of similar structure which, instead of the morpholino group, contain a diethanolamino group, a methylamino group, an anilino group or a 2-methoxyethylamino 15 group. Brighteners of the substituted diphenyl styryl type, for example alkali metal salts of 4,4'-bis-(2-sulfostyryl)-diphenyl, 4,4'-bis-(4-chloro-3-sulfostyryl)-diphenyl or 4-(4-chlorostyryl)-4'-(2-sulfostyryl)-diphenyl, may also be present. Mixtures of the brighteners mentioned above may also be used. Uniformly white granules are obtained if, in addition to the usual 20 brighteners in the usual quantities, for example between 0.1 and 0.5% by weight and preferably between 0.1 and 0.3% by weight, the detergents also contain small quantities, for example 10^{-6} to 10^{-3} % by weight and preferably around 10^{-5} % by weight, of a blue dye. A particularly preferred dye is Tinolux® (a product of Ciba-Geigy).

25 Suitable **soil repellents** are substances which preferably contain ethylene terephthalate and/or polyethylene glycol terephthalate groups, the molar ratio of ethylene terephthalate to polyethylene glycol terephthalate being in the range from 50:50 to 90:10. The molecular weight of the linking polyethylene glycol units is more particularly in the range from 750 to 30 5,000, i.e. the degree of ethoxylation of the polymers containing poly-

ethylene glycol groups may be about 15 to 100. The polymers are distinguished by an average molecular weight of about 5,000 to 200,000 and may have a block structure, but preferably have a random structure. Preferred polymers are those with molar ethylene terephthalate:

- 5 polyethylene glycol terephthalate ratios of about 65:35 to about 90:10 and preferably in the range from about 70:30 to 80:20. Other preferred polymers are those which contain linking polyethylene glycol units with a molecular weight of 750 to 5,000 and preferably in the range from 1,000 to about 3,000 and which have a molecular weight of the polymer of about
- 10 10,000 to about 50,000. Examples of commercially available polymers are the products Milease® T (ICI) or Repelotex® SRP 3 (Rhône-Poulenc).

Where the detergents are used in washing machines, it can be of advantage to add conventional **foam inhibitors** to them. Suitable foam inhibitors are, for example, soaps of natural or synthetic origin which have a high percentage of C₁₈₋₂₄ fatty acids. Suitable non-surface-active foam inhibitors are, for example, organopolysiloxanes and mixtures thereof with microfine, optionally silanized silica and paraffins, waxes, microcrystalline waxes and mixtures thereof with silanized silica or bis-stearyl ethylenediamide. Mixtures of various foam inhibitors, for example mixtures of silicones, paraffins or waxes, may also be used with advantage. The foam inhibitors, more particularly silicone- or paraffin-containing foam inhibitors, are preferably fixed to a granular water-soluble or water-dispersible carrier/support. Mixtures of paraffins and bis-stearyl ethylenediamides are particularly preferred.

- 25 The **pH** value of liquid detergents, more especially concentrated liquid detergents, is generally in the range from 7 to 10.5, preferably in the range from 7 to 9.5 and more preferably in the range from 7 to 8.5. Higher pH values, for example above 9, can be adjusted by using small quantities of sodium hydroxide or alkaline salts, such as sodium carbonate or sodium silicate. The liquid preparations generally have viscosities of 150 to 10,000
- 30

mPas (Brookfield viscosimeter, spindle 1, 20 r.p.m., 20°C). The substantially water-free detergents preferably have viscosities of 150 to 5,000 mPas. The viscosity of aqueous detergents is preferably below 2,000 mPas and, more particularly, in the range from 150 to 1,000 mPas.

5

Production of solid compositions

The bulk density of the solid compositions is generally in the range from 300 to 1,200 g/l and more particularly in the range from 500 to 1,100 g/l. They may be produced by any of the known methods, such as mixing, 10 spray drying, granulation and extrusion. Processes in which several components, for example spray-dried components and granulated and/or extruded components, are mixed with one another are particularly suitable. The spray dried or granulated components may also be subsequently treated, for example with nonionic surfactants, more particularly 15 ethoxylated fatty alcohols, by any of the usual methods. In granulation and extrusion processes in particular, any anionic surfactants present are preferably used in the form of a spray dried, granulated or extruded compound either as an added component in the process or as an additive to other granules. The preferred relatively heavy granules with bulk 20 densities above 600 g/l in particular preferably contain components which improve the dispensing behavior and/or the dissolving behavior of the granules. Alkoxylated fatty alcohols containing 12 to 80 moles of ethylene oxide per mole of alcohol, for example tallow fatty alcohol containing 14 EO, 30 EO or 40 EO, and polyethylene glycols with a relative molecular 25 weight of 200 to 12,000 and preferably in the range from 200 to 600, are advantageously used for this purpose.

It is also possible and, depending on the formulation, can be of advantage subsequently to add other individual ingredients of the detergent, for example citrate or citric acid or other polycarboxylates or 30 polycarboxylic acids, polymeric polycarboxylates, zeolite and/or layer

- silicates, which may optionally be crystalline, to spray-dried, granulated and/or extruded components optionally treated with nonionic surfactants and/or other ingredients which are liquid to wax-like at the processing temperature. A preferred process in this regard is one in which the surface
- 5 of ingredients of the detergent or the detergent as a whole is subsequently treated to reduce the tackiness of the granules rich in nonionic surfactants and/or to improve their solubility. Suitable surface modifiers are known from the prior art. Besides other suitable surface modifiers, fine-particle zeolites, silicas, amorphous silicates, fatty acids or fatty acid salts, for
- 10 example calcium stearate, but especially mixtures of zeolite and silicas or zeolite and calcium stearate, are particularly preferred.

Cosmetic and/or pharmaceutical preparations

- The branched, substantially unsaturated fatty alcohol sulfates according to the invention may be used for the production of cosmetic and/or pharmaceutical preparations, for example hair shampoos, hair lotions, foam baths, shower baths, creams, gels, lotions, alcoholic and aqueous/alcoholic solutions, emulsions, wax/fat compounds, stick preparations, powders or ointments. These preparations may also contain
- 15 mild surfactants, oil components, emulsifiers, superfatting agents, pearlizing waxes, consistency factors, thickeners, polymers, silicone compounds, fats, waxes, stabilizers, biogenic agents, deodorants, antiperspirants, antidandruff agents, film formers, swelling agents, UV protection factors, antioxidants, hydrotropes, preservatives, insect
- 20 repellents, self-tanning agents, solubilizers, perfume oils, dyes and the like as further auxiliaries and additives.

- Typical examples of suitable mild, i.e. particularly dermatologically compatible, **surfactants** are fatty alcohol polyglycol ether sulfates, monoglyceride sulfates, mono- and/or dialkyl sulfosuccinates, fatty acid
- 30 isethionates, fatty acid sarcosinates, fatty acid taurides, fatty acid

glutamates, α -olefin sulfonates, ether carboxylic acids, alkyl oligogluco-sides, fatty acid glucamides, alkylamidobetaines and/or protein fatty acid condensates, preferably based on wheat proteins.

- Suitable oil components are, for example, Guerbet alcohols based
- 5 on fatty alcohols containing 6 to 18 and preferably 8 to 10 carbon atoms, esters of linear C₆₋₂₂ fatty acids with linear C₆₋₂₂ fatty alcohols, esters of branched C₆₋₁₃ carboxylic acids with linear C₆₋₂₂ fatty alcohols such as, for example, myristyl myristate, myristyl palmitate, myristyl stearate, myristyl isostearate, myristyl oleate, myristyl behenate, myristyl erucate, cetyl
- 10 myristate, cetyl palmitate, cetyl stearate, cetyl isostearate, cetyl oleate, cetyl behenate, cetyl erucate, stearyl myristate, stearyl palmitate, stearyl stearate, stearyl isostearate, stearyl oleate, stearyl behenate, stearyl erucate, isostearyl myristate, isostearyl palmitate, isostearyl stearate, isostearyl isostearate, isostearyl oleate, isostearyl behenate, isostearyl
- 15 oleate, oleyl myristate, oleyl palmitate, oleyl stearate, oleyl isostearate, oleyl oleate, oleyl behenate, oleyl erucate, behenyl myristate, behenyl palmitate, behenyl stearate, behenyl isostearate, behenyl oleate, behenyl behenate, behenyl erucate, erucyl myristate, erucyl palmitate, erucyl stearate, erucyl isostearate, erucyl oleate, erucyl behenate and erucyl
- 20 erucate. Also suitable are esters of linear C₆₋₂₂ fatty acids with branched alcohols, more particularly 2-ethyl hexanol, esters of hydroxycarboxylic acids with linear or branched C₆₋₂₂ fatty alcohols, more especially Diethyl Malate, esters of linear and/or branched fatty acids with polyhydric alcohols (for example propylene glycol, dimer diol or trimer triol) and/or Guerbet
- 25 alcohols, triglycerides based on C₆₋₁₀ fatty acids, liquid mono-/di-/tri-glyceride mixtures based on C₆₋₁₈ fatty acids, esters of C₆₋₂₂ fatty alcohols and/or Guerbet alcohols with aromatic carboxylic acids, more particularly benzoic acid, esters of C₂₋₁₂ dicarboxylic acids with linear or branched alcohols containing 1 to 22 carbon atoms or polyols containing 2 to 10
- 30 carbon atoms and 2 to 6 hydroxyl groups, vegetable oils, branched primary

- alcohols, substituted cyclohexanes, linear and branched C₆-22 fatty alcohol carbonates, Guerbet carbonates, esters of benzoic acid with linear and/or branched C₆-22 alcohols (for example Finsolv® TN), linear or branched, symmetrical or nonsymmetrical dialkyl ethers containing 6 to 22 carbon atoms per alkyl group, ring opening products of epoxidized fatty acid esters with polyols, silicone oils and/or aliphatic or naphthenic hydrocarbons, for example squalane, squalene or dialkyl cyclohexanes.
- 5

Suitable **emulsifiers** are, for example, nonionic surfactants from at least one of the following groups:

- 10
- products of the addition of 2 to 30 moles of ethylene oxide and/or 0 to 5 moles of propylene oxide onto linear C₈-22 fatty alcohols, C₁₂-22 fatty acids and alkyl phenols containing 8 to 15 carbon atoms in the alkyl group and alkylamines containing 8 to 22 carbon atoms in the alkyl group;

15

 - alkyl and/or alkenyl oligoglycosides containing 8 to 22 carbon atoms in the alkyl group and ethoxylated analogs thereof;
 - products of the addition of 1 to 15 moles of ethylene oxide with castor oil and/or hydrogenated castor oil;

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 - products of the addition of 15 to 60 moles of ethylene oxide with castor oil and/or hydrogenated castor oil;
 - partial esters of glycerol and/or sorbitan with unsaturated, linear or saturated, branched fatty acids containing 12 to 22 carbon atoms and/or hydroxycarboxylic acids containing 3 to 18 carbon atoms and addition products thereof with 1 to 30 moles of ethylene oxide;

25

 - partial esters of polyglycerol (average degree of self-condensation 2 to 8), polyethylene glycol (molecular weight 400 to 5,000), trimethylolpropane, pentaerythritol, sugar alcohols (for example sorbitol), alkyl glucosides (for example methyl glucoside, butyl glucoside, lauryl glucoside) and polyglucosides (for example

30

cellulose) with saturated and/or unsaturated, linear or branched fatty acids containing 12 to 22 carbon atoms and/or hydroxycarboxylic acids containing 3 to 18 carbon atoms and addition products thereof with 1 to 30 moles of ethylene oxide;

- 5 ➤ mixed esters of pentaerythritol, fatty acids, citric acid and fatty alcohol according to DE 11 65 574 PS and/or mixed esters of fatty acids containing 6 to 22 carbon atoms, methyl glucose and polyols, preferably glycerol or polyglycerol,
- 10 ➤ mono-, di- and trialkyl phosphates and mono-, di- and/or tri-PEG- alkyl phosphates and salts thereof,
- wool wax alcohols,
- polysiloxane/polyalkyl/polyether copolymers and corresponding derivatives,
- polyalkylene glycols and
- 15 ➤ glycerol carbonate.

The addition products of ethylene oxide and/or propylene oxide with fatty alcohols, fatty acids, alkylphenols or with castor oil are known commercially available products. They are homolog mixtures of which the 20 average degree of alkoxylation corresponds to the ratio between the quantities of ethylene oxide and/or propylene oxide and substrate with which the addition reaction is carried out. C_{12/18} fatty acid monoesters and diesters of addition products of ethylene oxide with glycerol are known as refatting agents for cosmetic formulations from DE 20 24 051 PS.

25 **Alkyl and/or alkenyl oligoglycosides**, their production and their use are known from the prior art. They are produced in particular by reacting glucose or oligosaccharides with primary alcohols containing 8 to 18 carbon atoms. So far as the glucoside unit is concerned, both monoglycosides in which a cyclic sugar unit is attached to the fatty alcohol 30 by a glycoside bond and oligomeric glycosides with a degree of

oligomerization of preferably up to about 8 are suitable. The degree of oligomerization is a statistical mean value on which the homolog distribution typical of such technical products is based.

Typical examples of suitable **partial glycerides** are hydroxystearic

- 5 acid monoglyceride, hydroxystearic acid diglyceride, isostearic acid monoglyceride, isostearic acid diglyceride, oleic acid monoglyceride, oleic acid diglyceride, ricinoleic acid monoglyceride, ricinoleic acid diglyceride, linoleic acid monoglyceride, linoleic acid diglyceride, linolenic acid monoglyceride, linolenic acid diglyceride, erucic acid monoglyceride, erucic 10 acid diglyceride, tartaric acid monoglyceride, tartaric acid diglyceride, citric acid monoglyceride, citric acid diglyceride, malic acid monoglyceride, malic acid diglyceride and technical mixtures thereof which may still contain small quantities of triglyceride from the production process. Addition products of 1 to 30 and preferably 5 to 10 moles of ethylene oxide with the partial 15 glycerides mentioned are also suitable.

- Suitable **sorbitan esters** are sorbitan monoisostearate, sorbitan sesquoisostearate, sorbitan diisostearate, sorbitan triisostearate, sorbitan monooleate, sorbitan sesquioleate, sorbitan dioleate, sorbitan trioleate, sorbitan monoerucate, sorbitan sesquierucate, sorbitan dierucate, sorbitan 20 trierucate, sorbitan monoricinoleate, sorbitan sesquiricinoleate, sorbitan diricinoleate, sorbitan triricinoleate, sorbitan monohydroxystearate, sorbitan sesquihydroxystearate, sorbitan dihydroxystearate, sorbitan trihydroxystearate, sorbitan monotartrate, sorbitan sesquitartrate, sorbitan ditartrate, sorbitan tritartrate, sorbitan monocitrate, sorbitan sesquicitrate, sorbitan 25 dicitrate, sorbitan tricitrate, sorbitan monomaleate, sorbitan sesquimaleate, sorbitan dimaleate, sorbitan trimaleate and technical mixtures thereof. Addition products of 1 to 30 and preferably 5 to 10 moles of ethylene oxide with the sorbitan esters mentioned are also suitable.

Typical examples of suitable **polyglycerol esters** are Polyglyceryl-2

- 30 Dipolyhydroxystearate (Dehymuls® PGPH), Polyglycerin-3-Diisostearate

- (Lameform® TGI), Polyglyceryl-4 Isostearate (Isolan® GI 34), Polyglyceryl-3 Oleate, Diisostearoyl Polyglyceryl-3 Diisostearate (Isolan® PDI), Polyglyceryl-3 Methylglucose Distearate (Tego Care® 450), Polyglyceryl-3 Beeswax (Cera Bellina®), Polyglyceryl-4 Caprate (Polyglycerol Caprate 5 T2010/90), Polyglyceryl-3 Cetyl Ether (Chimexane® NL), Polyglyceryl-3 Distearate (Cremophor® GS 32) and Polyglyceryl Polyricinoleate (Admul® WOL 1403), Polyglyceryl Dimerate Isostearate and mixtures thereof.

Examples of other suitable **polyolesters** are the mono-, di- and triesters of trimethylol propane or pentaerythritol with lauric acid, cocofatty acid, tallow fatty acid, palmitic acid, stearic acid, oleic acid, behenic acid and the like optionally reacted with 1 to 30 moles of ethylene oxide.

Other suitable emulsifiers are **zwitterionic surfactants**. Zwitterionic surfactants are surface-active compounds which contain at least one quaternary ammonium group and at least one carboxylate and one sulfonate group in the molecule. Particularly suitable zwitterionic surfactants are the so-called betaines, such as the N-alkyl-N,N-dimethyl ammonium glycinate, for example cocoalkyl dimethyl ammonium glycinate, N-acylaminopropyl-N,N-dimethyl ammonium glycinate, for example cocoacylaminopropyl dimethyl ammonium glycinate, and 2-alkyl-15 3-carboxymethyl-3-hydroxyethyl imidazolines containing 8 to 18 carbon atoms in the alkyl or acyl group and cocoacylaminooethyl hydroxyethyl carboxymethyl glycinate. The fatty acid amide derivative known under the CTFA name of *Cocamidopropyl Betaine* is particularly preferred. Ampholytic surfactants are also suitable emulsifiers. Ampholytic surfactants are surface-active compounds which, in addition to a C_{8/18} alkyl or acyl group, contain at least one free amino group and at least one -COOH- or -SO₃H- group in the molecule and which are capable of forming inner salts. Examples of suitable ampholytic surfactants are N-alkyl glycines, N-alkyl propionic acids, N-alkylaminobutyric acids, N-alkyliminodipropionic 20 acids, N-hydroxyethyl-N-alkylamidopropyl glycines, N-alkyl taurines, N-alkyl 25 30

sarcosines, 2-alkylaminopropionic acids and alkylaminoacetic acids containing around 8 to 18 carbon atoms in the alkyl group. Particularly preferred ampholytic surfactants are N-cocoalkylaminopropionate, cocoacylaminoethyl aminopropionate and C_{12/18} acyl sarcosine.

5 Finally, other suitable emulsifiers are **cationic surfactants**, those of the esterquat type, preferably methyl-quaternized difatty acid triethanolamine ester salts, being particularly preferred.

10 **Superfattening agents** may be selected from such substances as, for example, lanolin and lecithin and also polyethoxylated or acylated lanolin and lecithin derivatives, polyol fatty acid esters, monoglycerides and fatty acid alkanolamides, the fatty acid alkanolamides also serving as foam stabilizers.

15 Suitable **pearlizing waxes** are, for example, alkylene glycol esters, especially ethylene glycol distearate; fatty acid alkanolamides, especially cocofatty acid diethanolamide; partial glycerides, especially stearic acid monoglyceride; esters of polybasic, optionally hydroxysubstituted carboxylic acids with fatty alcohols containing 6 to 22 carbon atoms, especially long-chain esters of tartaric acid; fatty compounds, such as for example fatty alcohols, fatty ketones, fatty aldehydes, fatty ethers and fatty 20 carbonates which contain in all at least 24 carbon atoms, especially laurone and distearylether; fatty acids, such as stearic acid, hydroxystearic acid or behenic acid, ring opening products of olefin epoxides containing 12 to 22 carbon atoms with fatty alcohols containing 12 to 22 carbon atoms and/or polyols containing 2 to 15 carbon atoms and 2 to 10 hydroxyl groups and 25 mixtures thereof.

20 The **consistency factors** mainly used are fatty alcohols or hydroxyfatty alcohols containing 12 to 22 and preferably 16 to 18 carbon atoms and also partial glycerides, fatty acids or hydroxyfatty acids. A combination of these substances with alkyl oligoglucosides and/or fatty acid 30 N-methyl glucamides of the same chain length and/or polyglycerol poly-12-

hydroxystearates is preferably used.

- Suitable **thickeners** are, for example, Aerosil types (hydrophilic silicas), polysaccharides, more especially xanthan gum, guar-guar, agar-agar, alginates and tyloses, carboxymethyl cellulose and hydroxyethyl cellulose, also relatively high molecular weight polyethylene glycol mono-esters and diesters of fatty acids, polyacrylates (for example Carbopol® [Goodrich] or Synthalens® [Sigma]), polyacrylamides, polyvinyl alcohol and polyvinyl pyrrolidone, surfactants such as, for example, ethoxylated fatty acid glycerides, esters of fatty acids with polyols, for example pentaerythritol or trimethylol propane, narrow-range fatty alcohol ethoxylates or alkyl oligoglucosides and electrolytes, such as sodium chloride and ammonium chloride.

- Suitable **cationic polymers** are, for example, cationic cellulose derivatives such as, for example, the quaternized hydroxyethyl cellulose obtainable from Amerchol under the name of Polymer JR 400®, cationic starch, copolymers of diallyl ammonium salts and acrylamides, quaternized vinyl pyrrolidone/vinyl imidazole polymers such as, for example, Luviquat® (BASF), condensation products of polyglycols and amines, quaternized collagen polypeptides such as, for example, Lauryldimonium Hydroxypropyl Hydrolyzed Collagen (Lamequat® L, Grünau), quaternized wheat polypeptides, polyethyleneimine, cationic silicone polymers such as, for example, amodimethicone, copolymers of adipic acid and dimethylamino-hydroxypropyl diethylenetriamine (Cartaretine®, Sandoz), copolymers of acrylic acid with dimethyl diallyl ammonium chloride (Merquat® 550, Chemviron), polyaminopolyamides as described, for example, in FR 2 252 840 A and crosslinked water-soluble polymers thereof, cationic chitin derivatives such as, for example, quaternized chitosan, optionally in micro-crystalline distribution, condensation products of dihaloalkyls, for example dibromobutane, with bis-dialkylamines, for example bis-dimethylamino-1,3-propane, cationic guar gum such as, for example, Jaguar®CBS,

Jaguar®C-17, Jaguar®C-16 of Celanese, quaternized ammonium salt polymers such as, for example, Mirapol® A-15, Mirapol® AD-1, Mirapol® AZ-1 of Miranol.

- Suitable anionic, zwitterionic, amphoteric and nonionic polymers are, for example, vinyl acetate/crotonic acid copolymers, vinyl pyrrolidone/vinyl acrylate copolymers, vinyl acetate/butyl maleate/isobornyl acrylate copolymers, methyl vinyl ether/maleic anhydride copolymers and esters thereof, uncrosslinked and polyol-crosslinked polyacrylic acids, acrylamidopropyl trimethylammonium chloride/acrylate copolymers, octylacrylamide/methyl methacrylate/tert.-butylaminoethyl methacrylate/2-hydroxypropyl methacrylate copolymers, polyvinyl pyrrolidone, vinyl pyrrolidone/vinyl acetate copolymers, vinyl pyrrolidone/dimethylaminoethyl methacrylate/vinyl caprolactam terpolymers and optionally derivatized cellulose ethers and silicones.
- Suitable silicone compounds are, for example, dimethyl polysiloxanes, methylphenyl polysiloxanes, cyclic silicones and amino-, fatty acid-, alcohol-, polyether-, epoxy-, fluorine-, glycoside- and/or alkyl-modified silicone compounds which may be both liquid and resin-like at room temperature. Other suitable silicone compounds are simethicones which are mixtures of dimethicones with an average chain length of 200 to 300 dimethylsiloxane units and hydrogenated silicates. A detailed overview of suitable volatile silicones can be found in Todd et al. in **Cosm. Toil.** 91, 27 (1976).
- Typical examples of fats are glycerides while suitable waxes are inter alia natural waxes such as, for example, candelilla wax, carnauba wax, Japan wax, espartograss wax, cork wax, guaruma wax, rice oil wax, sugar cane wax, ouricury wax, montan wax, beeswax, shellac wax, spermaceti, lanolin (wool wax), uropygial fat, ceresine, ozocerite (earth wax), petrolatum, paraffin waxes, microwaxes; chemically modified waxes (hard waxes) such as, for example, montan ester waxes, sasol waxes,

hydrogenated jojoba waxes and synthetic waxes such as, for example, polyalkylene waxes and polyethylene glycol waxes.

Metal salts of fatty acids such as, for example, magnesium, aluminium and/or zinc stearate or ricinoleate may be used as **stabilizers**.

- 5 In the context of the invention, **biogenetic agents** are, for example, tocopherol, tocopherol acetate, tocopherol palmitate, ascorbic acid, deoxyribonucleic acid, retinol, bisabolol, allantoin, phytantriol, panthenol, AHA acids, amino acids, ceramides, pseudoceramides, essential oils, plant extracts and vitamin complexes.
- 10 Cosmetic **deodorants** counteract, mask or eliminate body odors. Body odors are formed through the action of skin bacteria on apocrine perspiration which results in the formation of unpleasant-smelling degradation products. Accordingly, deodorants contain active principles which act as germ inhibitors, enzyme inhibitors, odor absorbers or odor
15 maskers.

- Basically, suitable **germ inhibitors** are any substances which act against gram-positive bacteria such as, for example, 4-hydroxybenzoic acid and salts and esters thereof, N-(4-chlorophenyl)-N'-(3,4-dichlorophenyl)-urea, 2,4,4'-trichloro-2'-hydroxydiphenylether (triclosan), 4-
20 chloro-3,5-dimethylphenol, 2,2'-methylene-bis-(6-bromo-4-chlorophenol), 3-methyl-4-(1-methylethyl)-phenol, 2-benzyl-4-chlorophenol, 3-(4-chlorophenoxy)-propane-1,2-diol, 3-iodo-2-propinyl butyl carbamate, chlorhexidine, 3,4,4'-trichlorocarbanilide (TTC), antibacterial perfumes, thymol, thyme oil, eugenol, nettle oil, menthol, mint oil, farnesol, phenoxyethanol, 25 glycerol monolaurate (GML), diglycerol monocaprate (DMC), salicylic acid-N-alkylamides such as, for example, salicylic acid-n-octyl amide or salicylic acid-n-decyl amide.

- Suitable **enzyme inhibitors** are, for example, esterase inhibitors. Esterase inhibitors are preferably trialkyl citrates, such as trimethyl citrate, 30 tripropyl citrate, triisopropyl citrate, tributyl citrate and, in particular, triethyl

citrate (Hydagen® CAT, Henkel KGaA, Düsseldorf, FRG). Esterase inhibitors inhibit enzyme activity and thus reduce odor formation. Other esterase inhibitors are sterol sulfates or phosphates such as, for example, lanosterol, cholesterol, campesterol, stigmasterol and sitosterol sulfate or phosphate, dicarboxylic acids and esters thereof, for example glutaric acid, glutaric acid monoethyl ester, glutaric acid diethyl ester, adipic acid, adipic acid monoethyl ester, adipic acid diethyl ester, malonic acid and malonic acid diethyl ester, hydroxycarboxylic acids and esters thereof, for example citric acid, malic acid, tartaric acid or tartaric acid diethyl ester, and zinc glycinate.

Suitable **odor absorbers** are substances which are capable of absorbing and largely retaining the odor-forming compounds. They reduce the partial pressure of the individual components and thus also reduce the rate at which they spread. An important requirement in this regard is that perfumes must remain unimpaired. Odor absorbers are not active against bacteria. They contain, for example, a complex zinc salt of ricinoleic acid or special perfumes of largely neutral odor known to the expert as "fixateurs" such as, for example, extracts of labdanum or styrax or certain abietic acid derivatives as their principal component. Odor maskers are perfumes or perfume oils which, besides their odor-masking function, impart their particular perfume note to the deodorants. Suitable perfume oils are, for example, mixtures of natural and synthetic perfumes. Natural perfumes include the extracts of blossoms, stems and leaves, fruits, fruit peel, roots, woods, herbs and grasses, needles and branches, resins and balsams. Animal raw materials, for example civet and beaver, may also be used. Typical synthetic perfume compounds are products of the ester, ether, aldehyde, ketone, alcohol and hydrocarbon type. Examples of perfume compounds of the ester type are benzyl acetate, p-tert.butyl cyclohexylacetate, linalyl acetate, phenyl ethyl acetate, linalyl benzoate, benzyl formate, allyl cyclohexyl propionate, styrylpropionate and benzyl

salicylate. Ethers include, for example, benzyl ethyl ether while aldehydes include, for example, the linear alkanals containing 8 to 18 carbon atoms, citral, citronellal, citronellyloxyacetaldehyde, cyclamen aldehyde, hydroxy-citronellal, lilial and bourgeonal. Examples of suitable ketones are the

5 ionones and methyl cedryl ketone. Suitable alcohols are anethol, citronellol, eugenol, isoeugenol, geraniol, linalool, phenylethyl alcohol and terpineol. The hydrocarbons mainly include the terpenes and balsams. However, it is preferred to use mixtures of different perfume compounds which, together, produce an agreeable fragrance. Other suitable perfume oils are essential

10 oils of relatively low volatility which are mostly used as aroma components. Examples are sage oil, camomile oil, clove oil, melissa oil, mint oil, cinnamon leaf oil, lime-blossom oil, juniper berry oil, vetiver oil, olibanum oil, galbanum oil, Iadanum oil and lavendin oil. The following are preferably used either individually or in the form of mixtures: bergamot oil,

15 dihydromyrcenol, lilial, lyral, citronellol, phenylethyl alcohol, α -hexylcinnamaldehyde, geraniol, benzyl acetone, cyclamen aldehyde, linalool, Boisambrene Forte, Ambroxan, indole, hedione, sandelice, citrus oil, mandarin oil, orange oil, allylamyl glycolate, cyclovertal, lavendin oil, clary oil, β -damascone, geranium oil bourbon, cyclohexyl salicylate,

20 Vertofix Coeur, Iso-E-Super, Fixolide NP, evernol, iraldein gamma, phenylacetic acid, geranyl acetate, benzyl acetate, rose oxide, romillat, irotyl and floramat.

25 **Antiperspirants** reduce perspiration and thus counteract underarm wetness and body odor by influencing the activity of the eccrine sweat glands. Aqueous or water-free antiperspirant formulations typically contain the following ingredients:

- 30 ➤ astringent active principles,
- oil components,
- nonionic emulsifiers,

- co-emulsifiers,
 - consistency factors,
 - auxiliaries in the form of, for example, thickeners or complexing agents and/or
- 5 ➤ nonaqueous solvents such as, for example, ethanol, propylene glycol and/or glycerol.

Suitable astringent active principles of antiperspirants are, above all, salts of aluminium, zirconium or zinc. Suitable antihydrotic agents of this 10 type are, for example, aluminium chloride, aluminium chlorohydrate, aluminium dichlorohydrate, aluminium sesquichlorohydrate and complex compounds thereof, for example with 1,2-propylene glycol, aluminium hydroxyallantoinate, aluminium chloride tartrate, aluminium zirconium trichlorohydrate, aluminium zirconium tetrachlorohydrate, aluminium 15 zirconium pentachlorohydrate and complex compounds thereof, for example with amino acids, such as glycine. Oil-soluble and water-soluble auxiliaries typically encountered in antiperspirants may also be present in relatively small amounts. Oil-soluble auxiliaries such as these include, for example,

- 20
- inflammation-inhibiting, skin-protecting or pleasant-smelling essential oils,
 - synthetic skin-protecting agents and/or
 - oil-soluble perfume oils.

- 25
- Typical water-soluble additives are, for example, preservatives, water-soluble perfumes, pH adjusters, for example buffer mixtures, water-soluble thickeners, for example water-soluble natural or synthetic polymers such as, for example, xanthan gum, hydroxyethyl cellulose, polyvinyl 30 pyrrolidone or high molecular weight polyethylene oxides.

Suitable **antidandruff agents** are climbazol, octopirox and zinc pyrithione.

- Standard **film formers** are, for example, chitosan, microcrystalline chitosan, quaternized chitosan, polyvinyl pyrrolidone, vinyl pyrrolidone/vinyl 5 acetate copolymers, polymers of the acrylic acid series, quaternary cellulose derivatives, collagen, hyaluronic acid and salts thereof and similar compounds.

- Suitable **swelling agents** for aqueous phases are montmorillonites, clay minerals, Pemulen and alkyl-modified Carbopol types (Goodrich). 10 Other suitable polymers and swelling agents can be found in R. Lochhead's review in **Cosm. Toil.** 108, 95 (1993).

- Examples of **UV protection factors** include organic substances (light filters) which are liquid or crystalline at room temperature and which are capable of absorbing ultraviolet radiation and of releasing the energy 15 absorbed in the form of longer-wave radiation, for example heat. UV-B filters can be oil-soluble or water-soluble. The following are examples of oil-soluble substances:

- 20 ➤ 3-benzylidene camphor or 3-benzylidene norcamphor and derivatives thereof, for example 3-(4-methylbenzylidene)-camphor, as described in **EP 0693471 B1**;
- 25 ➤ 4-aminobenzoic acid derivatives, preferably 4-(dimethylamino)-benzoic acid-2-ethylhexyl ester, 4-(dimethylamino)-benzoic acid-2-octyl ester and 4-(dimethylamino)-benzoic acid amyl ester;
- 30 ➤ esters of cinnamic acid, preferably 4-methoxycinnamic acid-2-ethylhexyl ester, 4-methoxycinnamic acid propyl ester, 4-methoxycinnamic acid isoamyl ester, 2-cyano-3,3-phenylcinnamic acid-2-ethylhexyl ester (Octocrylene);
- esters of salicylic acid, preferably salicylic acid-2-ethylhexyl ester, salicylic acid-4-isopropylbenzyl ester, salicylic acid homomenthyl ester;

- derivatives of benzophenone, preferably 2-hydroxy-4-methoxybenzophenone, 2-hydroxy-4-methoxy-4'-methylbenzophenone, 2,2'-dihydroxy-4-methoxybenzophenone;
- esters of benzalmalonic acid, preferably 4-methoxybenzalmalonic acid di-2-ethylhexyl ester;
- 5 ➤ triazine derivatives such as, for example, 2,4,6-trianilino-(p-carbo-2'-ethyl-1'-hexyloxy)-1,3,5-triazine and Octyl Triazone, as described in EP 0 818 450 A1, or Dioctyl Butamido Triazine (Uvasorb® HEB);
- 10 ➤ propane-1,3-diones such as, for example, 1-(4-tert.butylphenyl)-3-(4'-methoxyphenyl)-propane-1,3-dione;
- ketotricyclo(5.2.1.0)decane derivatives, as described in EP 0 694 521 B1.

Suitable water-soluble substances are

- 15 ➤ 2-phenylbenzimidazole-5-sulfonic acid and alkali metal, alkaline earth metal, ammonium, alkylammonium, alkanolammonium and glucammonium salts thereof;
- sulfonic acid derivatives of benzophenones, preferably 2-hydroxy-4-methoxybenzophenone-5-sulfonic acid and salts thereof;
- 20 ➤ sulfonic acid derivatives of 3-benzylidene camphor such as, for example, 4-(2-oxo-3-bornylidene)methyl)-benzene sulfonic acid and 2-methyl-5-(2-oxo-3-bornylidene)-sulfonic acid and salts thereof.

- 25 Typical UV-A filters are, in particular, derivatives of benzoyl methane such as, for example 1-(4'-tert.butylphenyl)-3-(4'-methoxyphenyl)-propane-1,3-dione, 4-tert-butyl-4'-methoxydibenzoylmethane (Parsol 1789), 1-phenyl-3-(4'-isopropylphenyl)-propane-1,3-dione and the eneamine compounds described in DE 19712033 A1 (BASF). The UV-A and UV-B filters may of course also be used in the form of mixtures. Besides the
- 30

soluble substances mentioned, insoluble pigments, i.e. finely dispersed metal oxides or salts, may also be used for this purpose. Examples of suitable metal oxides are, in particular, zinc oxide and titanium dioxide and also oxides of iron, zirconium, silicon, manganese, aluminium and cerium 5 and mixtures thereof. Silicates (talcum), barium sulfate and zinc stearate may be used as salts. The oxides and salts are used in the form of the pigments for skin-care and skin-protecting emulsions. The particles should have an average diameter of less than 100 nm, preferably from 5 to 50 nm and more preferably from 15 to 30 nm. They may be spherical in shape 10 although ellipsoidal particles or other non-spherical particles may also be used. The pigments may also be surface-treated, i.e. hydrophilicized or hydrophobicized. Typical examples are coated titanium dioxides such as, for example, Titandioxid T 805 (Degussa) or Eusolex® T2000 (Merck). Suitable hydrophobic coating materials are, above all, silicones and 15 particularly trialkoxyoctyl silanes or simethicones. So-called micro- or nanopigments are preferably used in sun protection products. Micronized zinc oxide is preferably used. Other suitable UV filters can be found in P. Finkel's review in **SÖFW-Journal 122, 543 (1996)**.

Besides the two above-mentioned groups of primary protection 20 factors, secondary protection factors of the **antioxidant** type may also be used. Secondary sun protection factors of the antioxidant type interrupt the photochemical reaction chain which is initiated when UV rays penetrate into the skin. Typical examples of suitable antioxidants are amino acids (for example glycine, histidine, tyrosine, tryptophane) and derivatives thereof, 25 imidazoles (for example urocanic acid) and derivatives thereof, peptides, such as D,L-carnosine, D-carnosine, L-carnosine and derivatives thereof (for example anserine), carotinoids, carotenes (for example α -carotene, β -carotene, lycopene) and derivatives thereof, chlorogenic acid and derivatives thereof, liponic acid and derivatives thereof (for example dihydroliponic acid), aurothioglucose, propylthiouracil and other thiols (for 30

example thioredoxine, glutathione, cysteine, cystine, cystamine and glycosyl, N-acetyl, methyl, ethyl, propyl, amyl, butyl and lauryl, palmitoyl, oleyl, γ -linoleyl, cholesteryl and glyceryl esters thereof) and their salts, dilaurylthiodipropionate, distearylthiodipropionate, thiodipropionic acid and derivatives thereof (esters, ethers, peptides, lipids, nucleotides, nucleosides and salts) and sulfoximine compounds (for example butionine sulfoximines, homocysteine sulfoximine, butionine sulfones, penta-, hexa- and hepta-thionine sulfoximine) in very small compatible dosages (for example pmole to μ mole/kg), also (metal) chelators (for example α -hydroxyfatty acids, palmitic acid, phytic acid, lactoferrine), α -hydroxy acids (for example citric acid, lactic acid, malic acid), humic acid, bile acid, bile extracts, bilirubin, biliverdin, EDTA, EGTA and derivatives thereof, unsaturated fatty acids and derivatives thereof (for example γ -linolenic acid, linoleic acid, oleic acid), folic acid and derivatives thereof, ubiquinone and ubiquinol and derivatives thereof, vitamin C and derivatives thereof (for example ascorbyl palmitate, Mg ascorbyl phosphate, ascorbyl acetate), tocopherols and derivatives (for example vitamin E acetate), vitamin A and derivatives (vitamin A palmitate) and coniferyl benzoate of benzoin resin, rutinic acid and derivatives thereof, α -glycosyl rutin, ferulic acid, furfurylidene glucitol, carnosine, butyl hydroxytoluene, butyl hydroxyanisole, nordihydroguaiac resin acid, nordihydroguaiaretic acid, trihydroxybutyrophene, uric acid and derivatives thereof, mannose and derivatives thereof, Superoxid-Dismutase, zinc and derivatives thereof (for example ZnO, ZnSO₄), selenium and derivatives thereof (for example selenium methionine), stilbenes and derivatives thereof (for example stilbene oxide, trans-stilbene oxide) and derivatives of these active substances suitable for the purposes of the invention (salts, esters, ethers, sugars, nucleotides, nucleosides, peptides and lipids).

In addition, **hydrotropes**, for example ethanol, isopropyl alcohol or polyols, may be used to improve flow behavior. Suitable polyols preferably

contain 2 to 15 carbon atoms and at least two hydroxyl groups. The polyols may contain other functional groups, more especially amino groups, or may be modified with nitrogen. Typical examples are

- 5 ➤ glycerol;
- alkylene glycols such as, for example, ethylene glycol, diethylene glycol, propylene glycol, butylene glycol, hexylene glycol and polyethylene glycols with an average molecular weight of 100 to 1000 dalton;
- technical oligoglycerol mixtures with a degree of self-condensation of 10 1.5 to 10 such as, for example, technical diglycerol mixtures with a diglycerol content of 40 to 50% by weight;
- methylol compounds such as, in particular, trimethylol ethane, trimethylol propane, trimethylol butane, pentaerythritol and dipentaerythritol;
- 15 ➤ lower alkyl glucosides, particularly those containing 1 to 8 carbon atoms in the alkyl group, for example methyl and butyl glucoside;
- sugar alcohols containing 5 to 12 carbon atoms, for example sorbitol or mannitol,
- sugars containing 5 to 12 carbon atoms, for example glucose or 20 sucrose;
- amino sugars, for example glucamine;
- dialcoholamines, such as diethanolamine or 2-aminopropane-1,3-diol.

25 Suitable **preservatives** are, for example, phenoxyethanol, formaldehyde solution, parabens, pentanediol or sorbic acid and the other classes of compounds listed in Appendix 6, Parts A and B of the Kosmetikverordnung ("Cosmetics Directive"). Suitable **insect repellents** are N,N-diethyl-m-toluamide, pentane-1,2-diol or Ethyl Butylacetyl-aminopropionate. A suitable **self-tanning agent** is dihydroxyacetone.

30 Suitable **perfume oils** are mixtures of natural and synthetic

perfumes. Natural perfumes include the extracts of blossoms (lily, lavender, rose, jasmine, neroli, ylang-ylang), stems and leaves (geranium, patchouli, petitgrain), fruits (anise, coriander, caraway, juniper), fruit peel (bergamot, lemon, orange), roots (nutmeg, angelica, celery, cardamon, 5 costus, iris, calamus), woods (pinewood, sandalwood, guaiac wood, cedarwood, rosewood), herbs and grasses (tarragon, lemon grass, sage, thyme), needles and branches (spruce, fir, pine, dwarf pine), resins and balsams (galbanum, elemi, benzoin, myrrh, olibanum, opopanax). Animal raw materials, for example civet and beaver, may also be used. Typical 10 synthetic perfume compounds are products of the ester, ether, aldehyde, ketone, alcohol and hydrocarbon type. Examples of perfume compounds of the ester type are benzyl acetate, phenoxyethyl isobutyrate, p-tert.butyl cyclohexylacetate, linalyl acetate, dimethyl benzyl carbinal acetate, phenyl ethyl acetate, linalyl benzoate, benzyl formate, ethylmethyl phenyl 15 glycinate, allyl cyclohexyl propionate, styrallyl propionate and benzyl salicylate. Ethers include, for example, benzyl ethyl ether while aldehydes include, for example, the linear alkanals containing 8 to 18 carbon atoms, citral, citronellal, citronellyloxyacetalddehyde, cyclamen aldehyde, hydroxycitronellal, lilial and bourgeonal. Examples of suitable ketones are 20 the ionones, α -isomethylionone and methyl cedryl ketone. Suitable alcohols are anethol, citronellol, eugenol, isoeugenol, geraniol, linalool, phenylethyl alcohol and terpineol. The hydrocarbons mainly include the terpenes and balsams. However, it is preferred to use mixtures of different perfume compounds which, together, produce an agreeable fragrance. 25 Other suitable perfume oils are essential oils of relatively low volatility which are mostly used as aroma components. Examples are sage oil, camomile oil, clove oil, melissa oil, mint oil, cinnamon leaf oil, lime-blossom oil, juniper berry oil, vetiver oil, olibanum oil, galbanum oil, ladanum oil and lavandin oil. The following are preferably used either individually or in the 30 form of mixtures: bergamot oil, dihydromyrcenol, lilial, lyral, citronellol,

phenylethyl alcohol, α -hexylcinnamaldehyde, geraniol, benzyl acetone, cyclamen aldehyde, linalool, Boisambrene Forte, Ambroxan, indole, hedione, sandelice, citrus oil, mandarin oil, orange oil, allylamyl glycolate, cyclovertal, lavandin oil, clary oil, β -damascone, geranium oil bourbon, 5 cyclohexyl salicylate, Vertofix Coeur, Iso-E-Super, Fixolide NP, evernyl, iraldein gamma, phenylacetic acid, geranyl acetate, benzyl acetate, rose oxide, romillat, irotyl and floramat.

10 Suitable dyes are any of the substances suitable and approved for cosmetic purposes as listed, for example, in the publication **"Kosmetische Färbemittel"** of the **Farbstoffkommission der Deutschen Forschungsgemeinschaft, Verlag Chemie, Weinheim, 1984, pages 81 to 106**. These dyes are normally used in concentrations of 0.001 to 0.1% by weight, based on the mixture as a whole.

15 The total percentage content of auxiliaries and additives may be from 1 to 50% by weight and is preferably from 5 to 40% by weight, based on the particular composition. The compositions may be produced by standard hot or cold processes and are preferably produced by the phase inversion temperature method.

20

Examples

25 **Example 1.** 23 kg of the monomer fatty acid Edenor® 935 (Henkel KGaA) were esterified with 20 kg of methanol for 2 h at 240°C/100 bar. After removal of the water/methanol mixture, the same quantity of fresh methanol was added and the procedure was repeated twice. The ester thus obtained had an acid value of 0.8. The methyl ester was 30 hydrogenated on a fixed-bed Zn/Cr catalyst with the double bond intact. The throughput of methyl ester was 0.5 unit by volume per hour, based on the total volume of the reactor. After removal of the methanol, the crude alcohol was distilled (3% first runnings, 90% main runnings, 6% residue).

The resulting alcohol had a hydroxyl value of 192, a saponification value of 0.9 and an iodine value of 74 (solidus point 25.8°C). 586 g (2 mol) of the isooleyl alcohol obtained were sulfated with sulfur trioxide in a molar ratio of 1:0.95 in a falling film reactor at a temperature of 25°C, the sulfur trioxide

- 5 being 3% diluted with nitrogen. Immediately after leaving the reactor, the reaction product was neutralized with aqueous sodium hydroxide solution and the neutralized product was post-hydrolyzed for 1 hour at pH 10. Analysis produced the following results:

10	dry residue	: 49.0% by weight
	detergent substance (Epton)	: 35.1% by weight
	unsulfated	: 13.0% by weight
	sodium sulfate	: 0.2% by weight

15

- Example 2.** Monomer fatty acid was substantially freed from straight-chain saturated fatty acids by crystallization from methanol/water (Emersol process). Around 20% by weight of fatty acid, predominantly palmitic and stearic acid, was removed in this way. The liquid fatty acid mixture 20 obtained after removal of the solvent by distillation had a titer of 5°C and was first converted into the methyl ester and then hydrogenated to the unsaturated fatty alcohol in the same way as in Example 1. The unsaturated fatty alcohol had a hydroxyl value of 191, a saponification value of 1.7 and an iodine value of 87 (solidus point 3.8°C). 586 g (2 mol) 25 of the isooleyl alcohol thus obtained were sulfated with sulfur trioxide in a molar ratio of 1:1.05 in a falling film reactor at a temperature of 30°C, the sulfur trioxide being 5% diluted with nitrogen. Immediately after leaving the reactor, the reaction product was neutralized with aqueous sodium hydroxide solution and the neutralized product was post-hydrolyzed for 1 30 hour at pH 10. Analysis produced the following results:

dry residue	: 49.8% by weight
detergent substance (Epton)	: 38.2% by weight
unsulfated	: 11.3% by weight
5 sodium sulfate	: 0.4% by weight

Tables 1 and 2 below shows a number of formulation examples ((1-6) light duty detergent, (7-9) heavy duty detergent, (10) toilet bar soap, (11-15) manual dishwashing detergent, (16,17) machine dishwashing 10 detergent, (18-20) cleaner, (21-24) hair rinse, (25-26) hair conditioner, (27-28) shower bath, (29) shower gel, (30) wash lotion, (31-34) "two-in-one" shower bath, (35-40) shampoo, (41-45) foam bath, (46) soft cream), (47,48) moisturizing emulsion, (49,50) night cream)

Table 1

Detergent preparations (water, preservative to 100% by weight)

Composition	1	2	3	4	5	6	7	8	9	10
Maranil® Paste A 55	-	23.6	-	19.0	-	-	11.0	7.0	-	-
Dodecybenzenesulfonate, sodium salt										
Fatty alcohol sulfate of Example 1	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	5.0	5.0
Sulfopon® 1218 W	5.5	-	14.5	-	-	-	-	3.0	7.0	30.0
Lauryl(stearyl) sulfate, sodium salt										
Texapon® N	-	-	-	-	-	9.6	-	-	-	-
Laurylalcohol + 2EO sulfate, sodium salt										
Dehyquart® AU 56	-	-	-	-	6.7	-	-	-	-	-
Methyl-quaternized dipalm oil fatty acid triethanolammonium ester, methosulfate										
Comperlan® 100	-	-	-	-	-	-	-	-	-	10.0
Cocofatty acid ethanolamide										
Dehydrol® O 8	-	-	-	-	-	30.0	-	-	-	-
Octanol + 8EO										
Dehydrol® LT5	-	-	-	-	-	-	10.0	-	6.0	-
Laurylstearyl alcohol + 5EO										
Dehydrol® LT 7	12.0	-	-	-	10.2	-	-	2.0	4.0	-
Laurylstearyl alcohol + 7EO										
Eumulgin® WO 7	-	-	10.0	4.0	-	-	-	-	-	-
Oleyl alcohol + 7 EO										
Eumulgin® RT 40	-	-	-	-	-	-	-	-	-	5.0
Castor oil + 40EO										
Glucopon® 600 CS UP	4.0	5.0	11.0	15.0	6.0	19.5	-	-	1.0	-
Lauryl glucoside										
Edenor® K 1218	6.0	-	4.0	-	-	12.0	-	-	-	-
Cocofatty acid										
Sokalan® CP5	-	-	-	5.0	-	-	-	4.0	5.5	-
Maleic acid/acrylic acid copolymer, sodium salt										
Sodium carbonate	-	2.0	-	7.0	-	-	13.0	7.0	10.0	-
Sodium sulfate	-	-	-	19.0	-	-	-	10.0	9.5	45.0
Sodium tripolyphosphate	-	20.0	-	-	-	-	-	-	-	-
Zeolite A	-	-	-	25.0	-	-	39.0	25.0	32.5	-
Amylase	-	-	-	-	-	-	0.2	-	0.2	-
Cellulase	-	-	-	-	-	-	0.2	-	-	-
Lipase	0.8	-	-	-	-	-	0.2	0.7	0.1	-
Protease	-	-	-	-	-	-	0.2	0.3	0.1	-
Glycerol	3.0	-	-	-	-	-	-	-	-	-
Propylene glycol	7.0	-	5.0	-	-	-	-	-	-	-
Ethanol	5.0	-	5.0	-	-	7.0	-	-	-	-
Protol® N	-	-	-	3.0	-	-	-	-	-	-
Sodium silicate										
Dehydran® 760	-	-	-	7.0	-	-	4.0	5.0	4.5	-
Silicon/Paraffin on carrier										
Polyvinyl pyrrolidone	-	-	-	-	-	-	0.3	1.0	0.8	-
Sodium hydroxide	5.5	-	5.0	-	-	7.6	-	-	-	-
Sodium hydrogen carbonate	-	-	-	-	-	-	7.0	9.0	5.5	-
Sodium citrate	0.1	-	-	-	0.1	-	4.0	12.0	4.0°	5.0

Table 1 continued

Detergent preparations (water, preservative to 100% by weight)

Composition	11	12	13	14	15	16	17	18	19	20
Maranil® Paste A 55	20.0	-	-	-	-	-	-	-	-	6.0
Dodecylbenzenesulfonate, sodium salt										
Fatty alcohol sulfate of Example 1	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Texapon® NSO	3.0	15.0	-	-	20.0	-	-	2.0	-	-
Lauryl alcohol+2EO sulfate, sodium salt										
Texapon® N70	-	-	9.0	-	-	-	-	-	-	-
Lauryl alcohol+2EO sulfate, sodium salt										
Texapon® LS 35	-	10.0	2.0	5.0	-	-	-	-	2.0	-
Lauryl alcohol+3.5EO sulfate, sodium salt										
Texapon® 842	5.0	3.5	-	-	-	-	-	-	-	-
Octyl sulfate, sodium salt										
Alkane sulfonate C13/17	-	-	-	12.5	18.0	-	-	-	-	-
Glucopon® 600 CS UP	3.0	2.0	15.0	4.0	-	-	-	1.0	-	-
Lauryl glucoside										
Dehydol® O10	-	-	-	-	8.0	-	-	-	-	-
Octanol+10EO										
Dehydol® 980	-	-	-	-	-	-	-	-	1.0	2.0
Laurylmystryl alcohol+1PO+6EO										
Dehypón® LS 54	-	-	-	-	-	15.0	-	-	-	-
Lauryl alcohol+5EO+4PO										
Dehypón® LS 104	-	-	-	-	-	-	10.0	-	-	-
Lauryl alcohol+10EO butyl ether										
Dehyton® K	-	3.5	5.0	-	-	-	-	-	-	-
Cocofatty acid amidopropyl betaine										
Sokalan® DCS	-	-	2.5	-	-	-	-	-	-	-
Succinic acid/glutaric acid/adipic acid mixture										
Ethanol	-	-	8.0	-	-	-	-	-	-	-
Isopropyl alcohol	-	-	-	-	-	-	-	-	-	3.0
Butyl diglycol	-	-	-	-	-	-	-	-	-	1.5
Cumene sulfonate, sodium salt	-	-	-	-	-	20.0	13.0	-	-	-
Sodium hydroxide	-	-	1.5	-	-	-	-	-	-	-
Ammonia	-	-	-	-	-	-	-	-	-	0.5
Hydrogen peroxide	-	-	-	-	-	-	-	5.0	5.0	-
Turpinal® 4NL	-	-	-	-	-	-	-	0.3	0.3	-
Tetrasodium edetionate										
Citric acid	0.2	0.2	0.2	0.2	0.2	-	5.0	-	-	-

Table 2

Cosmetic preparations (water, preservative to 100% by weight)

Composition (INCI)	21	22	23	24	25	26	27	28	29	30
Texapon® NSO	-	-	-	-	-	-	26.0	26.0	15.0	-
Sodium Laureth Sulfate										
Fatty alcohol sulfate of Example 1	1.0	1.0	1.0	1.0	1.0	1.0	12.0	12.0	10.0	1.0
Texapon® SB 3	-	-	-	-	-	-	-	-	10.0	-
Disodium Laureth Sulfosuccinate										
Plantacare® 818	-	-	-	-	-	-	7.0	7.0	6.0	-
Coco Glucosides										
Plantacare® PS 10	-	-	-	-	-	-	-	-	-	12.0
Sodium Laureth Sulfate (and) Coco Glucosides										
Dehyton® PK 45	-	-	-	-	-	-	-	-	10.0	-
Cocamidopropyl Betaine										
Dehyquart® A	2.0	2.0	2.0	2.0	4.0	4.0	-	-	-	-
Cetrimonium Chloride										
Dehyquart L® 80	1.2	1.2	1.2	1.2	0.6	0.6	-	-	-	-
Dococoylmethylethoxymonium Methosulfate (and) Propyleneglycol										
Eumulgin® B2	0.8	0.8	-	0.8	-	1.0	-	-	-	-
Ceteareth-20										
Eumulgin® VL 75	-	-	0.8	-	0.8	-	-	-	-	-
Lauryl Glucoside (and) Polyglyceryl-2										
Polyhydroxystearate (and) Glycerin										
Lanette® O	2.5	2.5	2.5	2.5	3.0	2.5	-	-	-	-
Cetearyl Alcohol										
Cutina® GMS	0.5	0.5	0.5	0.5	0.5	1.0	-	-	-	-
Glyceryl Stearate										
Cetiol® HE	1.0	-	-	-	-	-	-	-	1.0	-
PEG-7 Glyceryl Cocoate										
Cetiol® PGL	-	1.0	-	-	1.0	-	-	-	-	-
Hexyldecanol (and) Hexyldecyll laurate										
Eutanol® G	-	-	1.0	1.0	-	1.0	-	-	-	-
Octylidodecanol										
Nutrilan® Keratin W	-	-	-	2.0	-	-	-	-	-	-
Hydrolyzed Keratin										
Lamesoft® LMG	-	-	-	-	-	-	3.0	2.0	4.0	-
Glyceryl Laurate (and) Potassium Cocoyl										
Hydrolyzed Collagen										
Euperlan® PK 3000 AM	-	-	-	-	-	-	-	3.0	5.0	5.0
Glycol Distearate (and) Laureth-4 (and)										
Cocamidopropyl Betaine										
Generol® 122 N	-	-	-	-	1.0	1.0	-	-	-	-
Soja Sterol										
Highcareen® GS	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Betaglucan										
Hydagen® CMF	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Chitosan										
Copherol® 12250	-	-	0.1	0.1	-	-	-	-	-	-
Tocopherol Acetate										
Arlypon® F	-	-	-	-	-	-	3.0	3.0	1.0	-
Laureth-2										
Sodium Chloride	-	-	-	-	-	-	-	1.5	-	1.5

Table 2 continued

Cosmetic preparations (water, preservative to 100% by weight)

Composition (INCI)	31	32	33	34	35	36	37	38	39	40
Texapon® NSO	15.0	15.0	10.0	-	20.0	8.0	-	-	-	-
Sodium Laureth Sulfate										
Texapon® K 14 S	-	-	-	-	-	-	-	-	8.0	15.0
Sodium Myreth Sulfate										
Fatty alcohol sulfate of Example 1	5.0	5.0	2.4	1.0	5.0	3.0	1.0	1.0	3.0	8.0
Texapon® SB 3	-	-	-	-	-	7.0	-	-	-	-
Disodium Laureth Sulfosuccinate										
Plantacare® 818	5.0	5.0	4.0	-	-	-	-	-	6.0	4.0
Coco Glucosides										
Plantacare® 2000	-	-	-	-	5.0	4.0	-	-	-	-
Decyl Glucoside										
Plantacare® PS 10	-	-	-	40.0	-	-	16.0	17.0	-	-
Sodium Laureth Sulfate (and) Coco Glucosides										
Dehyton® PK 45	20.0	20.0	-	-	8.0	-	-	-	-	7.0
Cocamidopropyl Betaine										
Eumulgin® B1	-	-	-	-	1.0	-	-	-	-	-
Ceteareth-12										
Eumulgin® B2	-	-	-	1.0	-	-	-	-	-	-
Ceteareth-20										
Lameform® TGI	-	-	-	4.0	-	-	-	-	-	-
Polyglyceryl-3 Isostearate										
Dehytum® PGPH	-	-	-	1.0	-	-	-	-	-	-
Polyglyceryl-2 Dipolyhydroxystearate										
Monomuls® 90-L 12	-	-	-	-	-	-	-	-	1.0	1.0
Glyceryl Laurate										
Cetiol® HE	-	0.2	-	-	-	-	-	-	-	-
PEG-7 Glyceryl Cocoate										
Eutanol® G	-	-	-	1.0	-	-	-	-	-	-
Octyldodecanol										
Nutriian® Keratin W	-	-	-	-	-	-	-	-	2.0	2.0
Hydrolyzed Keratin										
Nutriian® I	1.0	-	-	-	-	2.0	-	2.0	-	-
Hydrolyzed Collagen										
Lamesoft® LMG	-	-	-	-	-	-	-	-	1.0	5.0
Glyceryl Laurate (and) Potassium Cocoyl Hydrolyzed Collagen										
Gluadin® WK	1.0	1.5	4.0	1.0	3.0	1.0	2.0	2.0	2.0	-
Sodium Cocoyl Hydrolyzed Wheat Protein										
Euperian® PK 3000 AM	5.0	3.0	4.0	-	-	-	-	3.0	3.0	-
Glycol Distearate (and) Laureth-4 (and) Cocamidopropyl Betaine										
Panthenol	-	-	1.0	-	-	-	-	-	-	-
Arlipon® F	2.6	1.6	-	1.0	1.5	-	-	-	-	-
Laureth-2										
Highcareen® GS	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Betaglucan										
Hydagen® CMF	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Chitosan										
Sodium Chloride	-	-	-	-	-	1.6	2.0	2.2	-	3.0
Glycerin (86% by weight)	-	5.0	-	-	-	-	-	1.0	3.0	-

Table 2 continued

Cosmetic preparations (water, preservative to 100 % by weight)

Composition (INCI)	41	42	43	44	45	46	47	48	49	50
Texapon® NSO	-	22.0	22.0	-	20.0	-	-	-	-	-
Sodium Laureth Sulfate										
Fatty alcohol sulfate of Example 2	1.0	8.0	8.0	1.0	5.0	1.0	1.0	1.0	1.0	1.0
Plantacare® 818	-	10.0	-	-	20.0	-	-	-	-	-
Coco Glucosides										
Plantacare® PS 10	22.0	-	5.0	22.0	-	-	-	-	-	-
Sodium Laureth Sulfate (and) Coco Glucosides										
Dehyton® PK 45	15.0	10.0	15.0	15.0	20.0	-	-	-	-	-
Cocamidopropyl Betaine										
Emulgade® SE	-	-	-	-	-	5.0	5.0	4.0	-	-
Glyceryl Stearate (and) Ceteareth 12/20 (and) Cetearyl Alcohol (and) Cetyl Palmitate										
Lameform® TGI	-	-	-	-	-	-	-	-	4.0	-
Polyglyceryl-3 Isostearate										
Dehymuls® PGPH	-	-	-	-	-	-	-	-	-	4.0
Polyglyceryl-2 Dipolyhydroxystearate										
Monomuls® 90-O 18	-	-	-	-	-	-	-	-	2.0	-
Glyceryl Oleate										
Cetiol® HE	2.0	-	-	2.0	5.0	-	-	-	-	2.0
PEG-7 Glyceryl Cocoate										
Cetiol® OE	-	-	-	-	-	-	-	-	5.0	6.0
Dicaprylyl Ether										
Cetiol® PGL	-	-	-	-	-	-	-	3.0	10.0	9.0
Hexyldecanol (and) Hexyldecyll Laurate										
Cetiol® SN	-	-	-	-	-	3.0	3.0	-	-	-
Cetearyl Isononanoate										
Cetiol® V	-	-	-	-	-	3.0	3.0	-	-	-
Decyl Oleate										
Myritol® 318	-	-	-	-	-	-	-	3.0	5.0	5.0
Coco Caprylate Caprate										
Bees Wax	-	-	-	-	-	-	-	-	7.0	5.0
Nutrilan® Elastin E20	-	-	-	-	-	2.0	-	-	-	-
Hydrolyzed Elastin										
Nutrilan® I-50	-	-	-	-	2.0	-	2.0	-	-	-
Hydrolyzed Collagen										
Gluadin® AGP	0.5	0.5	0.5	-	-	-	-	0.5	-	-
Hydrolyzed Wheat Gluten										
Gluadin® WK	2.0	2.0	2.0	2.0	5.0	-	-	-	0.5	0.5
Sodium Cocoyl Hydrolyzed Wheat Protein										
Euperlan® PK 3000 AM	5.0	-	-	5.0	-	-	-	-	-	-
Glycol Distearate (and) Laureth-4 (and) Cocamidopropyl Betaine										
Highcareen® GS	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Betaglucan										
Hydagen® CMF	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Chitosan										
Magnesium Sulfate Hepta Hydrate	-	-	-	-	-	-	-	-	1.0	1.0
Glycerin (95% by weight)	-	-	-	-	-	3.0	3.0	5.0	5.0	3.0

CLAIMS

1. Branched, substantially unsaturated fatty alcohol sulfates which are obtainable by
 - (a) dimerizing unsaturated C₁₆₋₂₂ fatty acids in known manner,
 - 5 (b) removing the monomer fraction accumulating in the dimerization step,
 - (c) converting the branched, substantially unsaturated fatty acids present in this fraction into the corresponding fatty acid methyl esters,
 - (d) hydrogenating the branched, substantially unsaturated fatty acid methyl esters with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols and
 - 10 (e) sulfating and neutralizing the branched, substantially unsaturated fatty alcohols in known manner with the double bond intact.
2. A process for the production of branched, substantially unsaturated fatty alcohol ether sulfates in which
 - (a) unsaturated C₁₆₋₂₂ fatty acids are dimerized in known manner,
 - (b) the monomer fraction accumulating in the dimerization step is removed,
 - 20 (c) the branched, substantially unsaturated fatty acids present in this fraction are converted into the corresponding fatty acid methyl esters,
 - (d) the branched, substantially unsaturated fatty acid methyl esters are hydrogenated with the double bonds intact to form the corresponding branched, substantially unsaturated fatty alcohols which are then
 - 25 (e) sulfated and neutralized in known manner.
3. A process as claimed in claim 2, characterized in that the monomer fraction accumulating in the dimerization step is first subjected to fractional crystallization and the liquid phase obtained is esterified, optionally after distillation.
- 30 4. A process as claimed in claims 2 and/or 3, characterized in that the

methyl esters and/or the fatty alcohols are subjected to distillation and/or fractional crystallization.

5. A process as claimed in at least one of claims 2 to 4, characterized in that the branched, substantially unsaturated fatty alcohols are sulfated with sulfur trioxide or chlorosulfonic acid.
6. A process as claimed in at least one of claims 2 to 5, characterized in that the branched, substantially unsaturated fatty alcohols are sulfated in a reactor operating on the falling-film principle.
7. A process as claimed in at least one of claims 2 to 6, characterized in that the branched, substantially unsaturated fatty alcohols are sulfated at temperatures of 25 to 90°C.
8. A process as claimed in at least one of claims 2 to 7, characterized in that the branched, substantially unsaturated fatty alcohols are sulfated in a molar ratio of fatty alcohol to sulfating agent of 1:0.95 to 1:1.8.
- 15 9. The use of the branched substantially unsaturated fatty alcohol sulfates claimed in claim 1 for the production of laundry detergents, dishwashing detergents, cleaners and softeners.
10. The use of the branched, substantially unsaturated fatty alcohol sulfates claimed in claim 1 for the production of cosmetic and/or pharmaceutical preparations.

(12) NACH DEM VERTRAG ÜBER DIE INTERNATIONALE ZUSAMMENARBEIT AUF DEM GEBIET DES PATENTWESENS (PCT) VERÖFFENTLICHTE INTERNATIONALE ANMELDUNG

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— Ohne internationalen Recherchenbericht und erneut zu
veröffentlichen nach Erhalt des Berichts.(71) Anmelder (für alle Bestimmungsstaaten mit Ausnahme
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Abkürzungen wird auf die Erklärungen ("Guidance Notes on
Codes and Abbreviations") am Anfang jeder regulären Ausgabe
der PCT-Gazette verwiesen.(72) Erfinder; und
(75) Erfinder/Anmelder (nur für US): WESTFECHTEL,

(54) Title: BRANCHED, SUBSTANTIALLY UNSATURATED FATTY ALCOHOL SULFATES

(54) Bezeichnung: VERZWEIGTE, WEITGEHEND UNGESÄTTIGTE FETTALKOHOLSULFATE

WO 01/14325 A2

(57) Abstract: The invention relates to branched, substantially unsaturated fatty alcohol sulfates that are obtained by a) dimerization of unsaturated fatty acids with 16 to 22 carbon atoms in a manner known per se, b) separation of the monomer fraction resulting from said dimerization, c) conversion of the branched, substantially unsaturated fatty acids contained in said fraction to the corresponding fatty acid methyl esters, d) hydration of the branched, substantially unsaturated fatty acid methyl esters to the corresponding branched, substantially unsaturated fatty alcohols, thereby obtaining double bonds, and e) sulfation and neutralization of said fatty alcohols in a manner known per se.

(57) Zusammenfassung: Vorgeschlagen werden verzweigte, weitgehend ungesättigte Fettsäurealkoholsulfate, die man erhält, indem man (a) ungesättigte Fettsäuren mit 16 bis 22 Kohlenstoffatomen in an sich bekannter Weise dimerisiert, (b) die bei der Dimerisierung anfallende Monomerfraktion abtrennt, (c) die in dieser Fraktion enthaltenen verzweigten, weitgehend ungesättigten Fettsäuren in die entsprechenden Fettsäuremethylester überführt, (d) die verzweigten, weitgehend ungesättigten Fettsäuremethylester unter Erhalt der Doppelbindungen zu den entsprechenden verzweigten, weitgehend ungesättigten Fettsäurealkoholen hydriert, und diese (e) in an sich bekannte Weise sulfatiert und neutralisiert.

"Express Mail" mailing label number EL541614378US.Type a plus sign (+) inside this box -

0010PTPO Rev. 6/95	U.S. Department of Commerce Patent and Trademark Office	Attorney Docket Number	C 2037 PCT/US
		First Named Inventor	HUEBNER, Norbert
COMPLETE IF KNOWN			
		Application Number	10/049,960
		Filing Date	06/19/2002
		Group Art Unit	
		Examiner Name	

Declaration Submitted with Initial Filing OR Declaration Submitted after Initial Filing

As a below named inventor, I hereby declare that:
My residence, post office address, and citizenship are as stated below next to my name.
I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

BRANCHED, SUBSTANTIALLY UNSATURATED FATTY ALCOHOL SULFATES

the specification of which	(Title of the Invention)		
<input type="checkbox"/> is attached hereto			
OR			
<input checked="" type="checkbox"/> was filed on (MM/DD/YYYY)	08/11/2000	as United States Application Number or PCT International	
Application Number	PCT/EP00/07849	and was amended on (MM/DD/YYYY)	(If applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37 Code of Federal Regulations, § 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code §119(e)-(6) or §365(b) of any foreign application(s) for patent or inventor's certificate, or §365(e) of any PCT International application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or of any PCT International application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached? YES	Certified Copy Attached? NO
199 39 565.9	DE	08/20/1999	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	<input type="checkbox"/>

Additional foreign application numbers are listed on a supplemental priority sheet attached hereto:

I hereby claim the benefit under Title 35, United States Code §119(e) of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (MM/DD/YYYY)	Additional provisional application numbers are listed on a supplemental priority sheet attached hereto.
		<input type="checkbox"/> application numbers are listed on a supplemental priority sheet attached hereto.

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DECLARATION

Page 2

I hereby claim the benefit under Title 35, United States Code §120 of any United States application(s), or §365(c) of any PCT International application designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code §112.1 acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations §1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

U.S. Parent Application Number	PCT Parent Number	Parent Filing Date (MM/DD/YYYY)	Parent Patent Number (if applicable)
	PCT/EP00/07849	08/11/2000	

Additional U.S. or PCT International application numbers are listed on a supplemental priority sheet attached hereto.

As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

<input type="checkbox"/> Firm Name OR <input checked="" type="checkbox"/> List Attorney(s) and/or agent(s) name and registration number below:	Customer Number or label		
Name	Registration Number	Name	Registration Number
John E. Drach Steven J. Trzaska	32,891 36,296	Aaron R. Ettelman Henry E. Millson, Jr.	42,516 18,980

Additional attorney(s) and/or agent(s) named on a supplemental sheet attached hereto.

Please direct all correspondence to:		<input checked="" type="checkbox"/> Customer Number or label	23657	OR	<input checked="" type="checkbox"/> Fill in correspondence address below
Name	John E. Drach				
Address					
Address					
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Country	Telephone		610-278-4925	Fax	610-278-4971

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Name of Sole or First Inventor:		<input type="checkbox"/> A petition has been filed for this unsigned				
Given Name	Norbert		Middle Initial	Family Name	Huebner	Suffix e.g. Jr.
Inventor's Signature	<i>Norbert Huebner</i>				Date	06.02.02
Residence: City	Langenfeld		State	Country	Germany	Citizenship
Post Office Address	AR Langenfeld 38					
Post Office Address						
City	40764 Langenfeld	State	Zip	Country	Germany	Applicant Authority

Additional inventors are being named on supplemental sheet(s) attached hereto

DECLARATION**ADDITIONAL INVENTOR(S)
Supplemental Sheet**Name of Additional Joint Inventor, if any: A petition has been filed for this unsigned inventor

Given Name	Hans-Christian		Middle Initial	Family Name	Raths	Suffix e.g. Jr.
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Inventor's Signature	<i>Hans - Christian Raths</i>			Date	21.02.02	
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Residence: City	Monheim	State		Country	Germany	Citizenship	Germany
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Post Office Address	Nelly-Sachs-Strasse 13						
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Post Office Address							
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City	40789 Monheim	State	Zip	Country	Germany	Applicant Authority
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Name of Additional Joint Inventor, if any: A petition has been filed for this unsigned inventor

Given Name	Alfred		Middle Initial	Family Name	Westfechtel	Suffix e.g. Jr.
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Inventor's Signature	<i>Alfred Westfechtel</i>			Date	27.2.02	
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Residence: City	Hilden	State		Country	Germany	Citizenship	Germany
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Post Office Address	Menzelweg 74						
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Post Office Address							
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City	40724 Hilden	State	Zip	Country	Germany	Applicant Authority
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Name of Additional Joint Inventor, if any: A petition has been filed for this unsigned inventor

Given Name			Middle Initial	Family Name				Suffix e.g. Jr.
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Inventor's Signature					Date		
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Residence: City		State		Country		Citizenship	
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Post Office Address							
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Post Office Address							
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City		State	Zip	Country		Applicant Authority
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Name of Additional Joint Inventor, if any: A petition has been filed for this unsigned inventor

Given Name			Middle Initial	Family Name				Suffix e.g. Jr.
------------	--	--	----------------	-------------	--	--	--	-----------------

Inventor's Signature					Date		
----------------------	--	--	--	--	------	--	--

Residence: City		State		Country		Citizenship	
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Post Office Address							
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Post Office Address							
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City		State	Zip	Country		Applicant Authority
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Additional inventors are being named on supplemental sheet(s) attached hereto